

=> s 11

SAMPLE SEARCH INITIATED 10:11:33 FILE 'MARPAT'
SAMPLE SCREEN SEARCH COMPLETED - 71 TO ITERATE
100.0% PROCESSED 71 ITERATIONS
SEARCH TIME: 00.00.09

2 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 915 TO 1925
PROJECTED ANSWERS: 2 TO 125

L2 2 SEA SSS SAM L1

=> s 11 full

FULL SEARCH INITIATED 10:11:48 FILE 'MARPAT'
FULL SCREEN SEARCH COMPLETED - 1426 TO ITERATE
85.0% PROCESSED 1212 ITERATIONS
98.5% PROCESSED 1405 ITERATIONS
100.0% PROCESSED 1426 ITERATIONS
SEARCH TIME: 00.00.46

74 ANSWERS
80 ANSWERS
80 ANSWERS

L3 80 SEA SSS FUL L1

=> fil caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	85.70	86.15

FILE 'CAPLUS' ENTERED AT 10:13:40 ON 13 NOV 1999
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FILE COVERS 1967 - 13 Nov 1999 VOL 131 ISS 21
FILE LAST UPDATED: 12 Nov 1999 (19991112/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

This file supports REGISTRY for direct browsing and searching of all substance data from the REGISTRY file. Enter HELP FIRST for more information.

=> s 13

L4 80 L3

=> s 14 and process

1206965 PROCESS
3 L4 AND PROCESS

=> s 14 and halogen?

93997 HALOGEN?
L6 2 L4 AND HALOGEN?

=> s 14 and thiazol?

29329 THIAZOL?
L7 33 L4 AND THIAZOL?

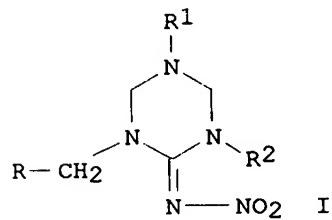
=> s 17 and process

1206965 PROCESS
L8 2 L7 AND PROCESS

=> d 15 1-3 all

L5 ANSWER 1 OF 3 CAPLUS COPYRIGHT 1999 ACS
AN 1998:668115 CAPLUS
DN 129:290052
TI Process for the preparation of nitroguanidine derivatives starting from 2-nitroiminohexahydro-1,3,5-triazines in the presence of ammonia, primary or secondary amines
IN Ebihara, Koichi; Ura, Daisuke; Miyamoto, Michihiko; Kaiho, Tatsuo
PA Mitsui Chemicals, Inc., Japan
SO Eur. Pat. Appl., 13 pp.
CODEN: EPXXDW
DT Patent
LA English
IC ICM C07D213-61
 ICS C07D277-32; C07D307-14
ICA C07D401-06; C07D417-06; C07D407-06
CC 27-6 (Heterocyclic Compounds (One Hetero Atom))

FAN.CNT 1		KIND	DATE	APPLICATION NO.	DATE
	PATENT NO.				
PI	EP 869120	A1	19981007	EP 1998-105850	19980331
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	CN 1197064	A	19981028	CN 1998-108267	19980331
	JP 11236381	A2	19990831	JP 1998-86842	19980331
PRAI	JP 1997-80178		19970331		
	JP 1997-82838		19970401		
	JP 1997-223813		19970820		
	JP 1997-258968		19970924		
	JP 1997-347934		19971217		
OS	MARPAT 129:290052				
GI					



AB Described is a process, as a substitute for hydrolysis, for prep. a nitroguanidine deriv., RCH₂NHC(:NNO₂)NHR₂ (R = 2-chloro-5-pyridyl, 2-chloro-5-thiazolyl, 2-, 3-tetrahydrofuryl, 2-methyl-4-tetrahydrofuryl; R₂ = Me, allyl), which comprises treating a triazine, I (R₁ = Me, benzyl, i-Pr, Et, t-Bu, cyclohexyl; R, R₂ = same as above), with NH₃, a primary amine or a secondary amine, or a salt thereof.

ST guanidine nitro prep; iminohexahydrotriazine amination; triazine hydro aminative ring cleavage

IT 136516-18-2 165253-14-5 168688-97-9 195986-55-1 213967-55-6
214149-49-2 214149-50-5

RL: RCT (Reactant)

(aminative ring cleavage of)

IT 131748-47-5P 131748-59-9P 165252-70-0P 165253-05-4P 214149-48-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

L5 ANSWER 2 OF 3 CAPLUS COPYRIGHT 1999 ACS

AN 1993:59590 CAPLUS

DN 118:59590

TI Heterocyclic amidine derivatives [(heteroaryl methyl)nitroguanidine derivatives] and their use as pesticides (insecticides and acaricides)

IN Kristiansen, Odd; Gsell, Laurenz; Maienfisch, Peter

PA Ciba-Geigy A.-G., Switz.

SO Eur. Pat. Appl., 63 pp.

CODEN: EPXXDW

DT Patent

LA German

IC ICM C07D213-40

ICS C07D277-32; C07D213-61; C07D213-89; A01N043-40; A01N043-78;
A01N047-42

CC 27-16 (Heterocyclic Compounds (One Hetero Atom))

Section cross-reference(s): 5

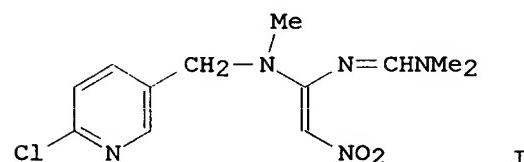
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 507736	A1	19921007	EP 1992-810225	19920326
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, PT				
	US 5223520	A	19930629	US 1992-858910	19920327
	CA 2064920	AA	19921005	CA 1992-2064920	19920402
	JP 05117237	A2	19930514	JP 1992-109199	19920402
	AU 9214036	A1	19921008	AU 1992-14036	19920403
	CN 1065456	A	19921021	CN 1992-102347	19920403
	HU 60721	A2	19921028	HU 1992-1140	19920403
	BR 9201197	A	19921201	BR 1992-1197	19920403
	ZA 9202450	A	19930405	ZA 1992-2450	19920403

PRAI CH 1991-1004255 19910404

OS CASREACT 118:59590; MARPAT 118:59590

GI

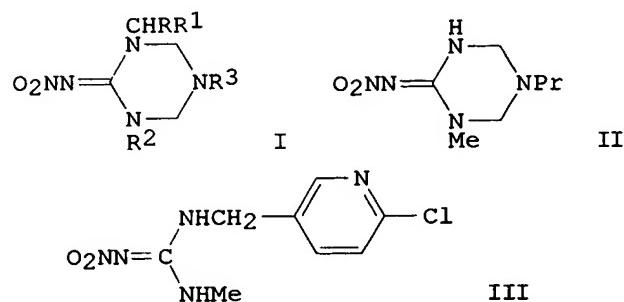


AB Some heterocyclic amidine derivs. and a process for their prepn. are claimed. The use of these compds. as pesticides (insecticides and acaricides) is claimed. Condensation of 1-amino-2-[(2-chloro-5-pyridyl)methylamino]-2-nitroethene with DMF di-Et acetal gave the acyclic amidine I. I had activity against Nilaparvata lugens, Nephotettix

ST cincticeps, Bemisia tabaci and Ctenocephalides felis (flea; systemic).
 heterocyclic amidine prepn pesticide; guanidine nitro heteroaryl methyl
 prepn insecticide acaricide; flea guanidine nitro heteroaryl methyl prepn
 IT Acaricides
 Insecticides
 Pesticides
 ((heteroaryl methyl)nitroguanidine derivs.)
 IT Ctenocephalides felis
 Flea
 (inhibition of, systemic, (heteroaryl methyl)nitroguanidine derivs.
 for)
 IT 85297-80-9 120738-75-2 131748-49-7 145369-00-2 145369-01-3
 RL: RCT (Reactant)
 (condensation reaction of, with DMF di-Et acetal)
 IT 1188-33-6, N,N-Dimethylformamide diethyl acetal
 RL: RCT (Reactant)
 (condensation reaction of, with
 [(chloropyridyl)methyl]methyl(nitro)gua
 nidine)
 IT 70258-18-3
 RL: RCT (Reactant)
 (condensation reaction of, with
 [(dimethylamino)methylene]methyl(nitro)
 guanidine)
 IT 145368-93-0 145368-94-1
 RL: RCT (Reactant)
 (condensation reaction of, with chloro(chloromethyl)pyridine)
 IT 14527-26-5
 RL: RCT (Reactant)
 (nitration of)
 IT 2986-25-6P 145368-95-2P 145368-96-3P 145368-97-4P 145368-98-5P
 145368-99-6P 145369-02-4P 145369-03-5P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of, as intermediate for (heteroaryl methyl)nitroguanidine
 (pesticide))
 IT 144930-61-0P 144930-63-2P 144930-75-6P 145368-66-7P 145368-67-8P
 145368-68-9P 145368-69-0P 145368-70-3P 145368-71-4P 145368-72-5P
 145368-73-6P 145368-74-7P 145368-75-8P 145368-76-9P 145368-77-0P
 145368-78-1P 145368-79-2P 145368-80-5P 145368-81-6P 145368-82-7P
 145368-83-8P 145368-84-9P 145368-85-0P 145368-86-1P 145368-87-2P
 145368-88-3P 145368-89-4P 145368-90-7P 145368-91-8P 145368-92-9P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except
 adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
 (Preparation); USES (Uses)
 (prepn. of, as pesticide)
 IT 765-30-0, Cyclopropylamine
 RL: RCT (Reactant)
 (reaction of, with S-methyl-N-nitroisothiourea)

L5 ANSWER 3 OF 3 CAPLUS COPYRIGHT 1999 ACS
 AN 1992:426353 CAPLUS
 DN 117:26353
 TI Process for the preparation of nitroguanidine derivatives
 IN Maienfisch, Peter; Kristiansen, Odd; Gsell, Laurenz
 PA Ciba-Geigy A.-G., Switz.
 SO Eur. Pat. Appl., 52 pp.
 CODEN: EPXXDW
 DT Patent
 LA German
 IC ICM C07D213-61
 ICS A01N043-40; C07D213-40; C07D277-32; A01N043-78
 CC 27-16 (Heterocyclic Compounds (One Hetero Atom))
 FAN.CNT 1
 PATENT NO. KIND DATE APPLICATION NO. DATE
 ----- -----

PI	EP 483062	A2	19920429	EP 1991-810795	19911015
	EP 483062	A3	19921028		
	EP 483062	B1	19960918		
	R: BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL				
	ES 2091899	T3	19961116	ES 1991-810795	19911015
	US 5245040	A	19930914	US 1991-777856	19911016
	IL 99801	A1	19960723	IL 1991-99801	19911018
	CA 2053954	AA	19920425	CA 1991-2053954	19911022
	JP 04330049	A2	19921118	JP 1991-303960	19911023
PRAI	CH 1990-3395		19901024		
OS	CASREACT 117:26353; MARPAT 117:26353				
GI					



AB O₂NN:C(NHR₂)NHCH₂R₁ (R = heterocyclic; R₁ = H, alkyl; R₂ = H, alkyl, aralkyl, cycloalkyl) were prep'd. by hydrolyzing triazines I [R₃ = (un)substituted alkyl, cycloalkyl, Ph, CH₂Ph]. Thus, 17.1 g O₂NN:C(NH₂)NHMe was treated with PrNH₂ and aq. CH₂OH to give 26.9 g triazine II. Treatment of 20.1 g II with 16.2 g 2-chloro-5-chloromethylpyridine to give 17.4 g I (R = 2-chloro-5-pyridyl, R₁, R₂ = H,

H, R₃ = Pr) which (4.25 g) was hydrolyzed with HCl-MeOH to give 2.51 g nitroguanidine III.

ST nitroguanidine pyridylmethyl thiazolylmethyl; pyridylmethylnitroguanidine; thiazolylmethylnitroguanidine

IT 141856-78-2P 141856-84-0P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. and hydrolysis of)

IT 141856-46-4P 141856-49-7P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. and reaction of, with chloro(chloromethyl)pyridine)

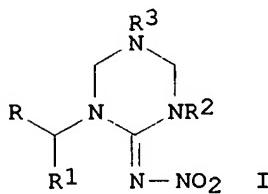
141856-65-7P	131748-47-5P	131748-50-0P	131748-53-3P	131748-56-6P
131748-59-9P	133258-61-4P	133258-65-8P	133258-66-9P	133258-70-5P
134323-25-4P	135018-04-1P	135018-10-9P	135018-11-0P	135018-14-3P
135018-15-4P	135018-16-5P	135018-17-6P	135018-18-7P	135018-19-8P
135018-20-1P	136516-16-0P	136516-17-1P	136516-18-2P	140920-89-4P
141856-47-5P	141856-48-6P	141856-50-0P	141856-51-1P	141856-52-2P
141856-53-3P	141856-54-4P	141856-55-5P	141856-56-6P	141856-57-7P
141856-58-8P	141856-59-9P	141856-60-2P	141856-61-3P	141856-62-4P
141856-63-5P	141856-64-6P	141856-65-7P	141856-66-8P	141856-67-9P
141856-68-0P	141856-69-1P	141856-70-4P	141856-71-5P	141856-72-6P
141856-73-7P	141856-74-8P	141856-75-9P	141856-76-0P	141856-77-1P
141856-79-3P	141856-80-6P	141856-81-7P	141856-82-8P	141856-83-9P
141856-85-1P	141856-86-2P	141856-87-3P	141856-88-4P	141856-89-5P
141856-90-8P	141856-91-9P	141856-92-0P	141856-93-1P	141856-94-2P
141856-95-3P	141856-96-4P	141856-97-5P	141856-98-6P	141856-99-7P
141857-00-3P	141857-01-4P	141857-02-5P	141857-03-6P	141857-04-7P
141857-05-8P	141857-06-9P	141857-07-0P	141857-08-1P	141857-09-2P
141857-10-5P	141857-11-6P	141857-12-7P	141857-13-8P	141857-14-9P
141857-15-0P	141857-16-1P	141857-17-2P	141857-18-3P	141857-19-4P

141857-20-7P 141857-21-8P 141857-22-9P 141857-23-0P 141857-24-1P
 141857-25-2P 141857-26-3P 141857-27-4P 141857-28-5P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of)
 IT 556-88-7, Nitroguanidine 4245-76-5
 RL: RCT (Reactant)
 (reaction of, with amines and formaldehyde)
 IT 62-53-3, Aniline, reactions 74-89-5, Methylamine, reactions 107-10-8,
 Propylamine, reactions
 RL: RCT (Reactant)
 (reaction of, with nitroguanidine and formaldehyde)
 IT 50-00-0, Formaldehyde, reactions
 RL: RCT (Reactant)
 (reaction of, with nitroguanidines and amines)
 IT 70258-18-3, 2-Chloro-5-chloromethylpyridine
 RL: RCT (Reactant)
 (reaction of, with nitroiminotriazacyclohexanes)

=> d 17 1-10 all

L7 ANSWER 1 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1999:136884 CAPLUS
 DN 130:182487
 TI Preparation of insecticidal and acaricidal 2-nitroguanidines
 IN Maienfisch, Peter
 PA Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft Mbh
 SO PCT Int. Appl., 31 pp.
 CODEN: PIXXD2
 DT Patent
 LA German
 IC ICM C07D213-61
 ICS C07D213-36; C07D277-32; C07D277-28; C07D213-89; C07D307-14;
 C07D261-10; C07D251-08; C07D403-04; C07D405-04; C07D417-12
 CC 28-21 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9909009	A1	19990225	WO 1998-EP5248	19980818
	W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
	AU 9893442	A1	19990308	AU 1998-93442	19980818
PRAI	CH 1997-1951		19970820		
	WO 1998-EP5248		19980818		
OS	MARPAT 130:182487				
GI					



AB 2-Nitroguanidines RR₁NHC(:NNO₂)NHR₂ [R = (un)substituted mono- or
bibcyclic heterocyclic; R₁ = H, alkyl; R₂ = H, alkyl, cycloalkyl,
(un)substituted benzyl, thiazolylmethyl, 3-pyridylmethyl] were
prep'd. by hydrolyzing a triazine I [R₃ = (un)substituted C₁₁-C₂₂ alkyl,
C₇-C₁₆-cycloalkyl, C₃-C₂₀ alkenyl, C₃-C₂₀ alkynyl, NH₂ aryl,
heterocyclic]. Thus, 1-methyl-2-nitroguanidine was cyclized with
octadecylamine and CH₂O and treated with 2-chloro-5-chloromethylpyridine
to give the triazine I [R = 2-chloro-5-pyridyl, R₁ = H, R₂ = Me, R₃ =
octadecyl] which was hydrolyzed with HCl in MeOH to give
MeNHC(:NNO₂)NHCH₂R.

ST nitroguanidine prep'n insecticide acaricide

IT Acaricides

Insecticides

(prepn. of insecticidal and acaricidal 2-nitroguanidines)

IT 131748-47-5P

RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of insecticidal and acaricidal 2-nitroguanidines)

IT 107-11-9, Allylamine 124-30-1, Octadecylamine 462-08-8,
3-Aminopyridine 4245-76-5 70258-18-3, 2-Chloro-5-chloromethylpyridine
105827-91-6, 2-Chloro-5-chloromethylthiazole 220620-28-0

RL: RCT (Reactant)

(prepn. of insecticidal and acaricidal 2-nitroguanidines)

IT 141856-64-6P 141857-04-7P 146270-52-2P 220620-11-1P 220620-12-2P
220620-13-3P 220620-14-4P 220620-15-5P 220620-16-6P 220620-17-7P
220620-18-8P 220620-19-9P 220620-20-2P 220620-21-3P 220620-22-4P
220620-23-5P 220620-24-6P 220620-25-7P 220620-26-8P 220620-27-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)

(prepn. of insecticidal and acaricidal 2-nitroguanidines)

L7 ANSWER 2 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1999:136883 CAPLUS

DN 130:182483

TI Preparation of insecticidal and acaricidal 2-nitroguanidines

IN Maienfisch, Peter

PA Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft
M.B.H.

SO PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DT Patent

LA German

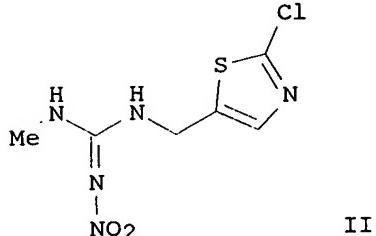
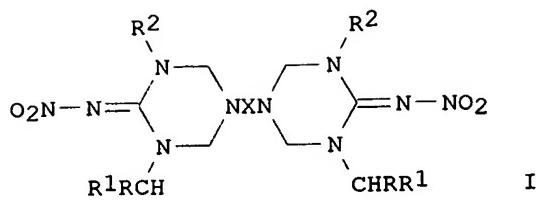
ICM C07D213-40

ICS C07D213-61; C07D213-89; C07D261-08; C07D277-28; C07D277-32;
C07D307-14; C07D251-08; C07D401-14; C07D405-14; C07D413-14;
C07D417-14

CC 28-12 (Heterocyclic Compounds (More Than One Hetero Atom)).
Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9909008	A1	19990225	WO 1998-EP5166	19980814
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	AU 9893423	A1	19990308	AU 1998-93423	19980814
PRAI	CH 1997-1934		19970818		
	WO 1998-EP5166		19980814		
OS	MARPAT 130:182483				
	GI				



AB 2-Nitroguanidines R2NHC(:NNO₂)NHCHRR₁ [R = (un) substituted heterocyclic; R₁ = H, alkyl; R₂ = H, alkyl, cycloalkyl, (un) substituted CH₂Ph, 3-pyridylmethyl, thienylmethyl] were obtained by hydrolyzing an alkylenebis(triazine) I [X = bond, alkylene, r₃ = CHRR₁]. Thus, 1-methyl-2-nitroguanidine was treated with ethylenediamine and CH₂O to give the triazine I [X = CH₂CH₂, R₃ = H] which was treated with 2-chloro-5-chloromethylthiazole to give I [R = 2-chloro-5-thiazolylmethyl, R₁ = H, R₂ = Me]. This latter compd. was treated with HCl in MeOH to give the nitroguanidine II.

ST nitroguanidine prep insecticide acaricide; nitroiminotriazine alkylenebis

IT Acaricides
Insecticide

(prepn. of insecticidal and acaricidal 2-nitroguanidines)

IT 131748-47-5P 131748-59-9P 131768-13-3P 135018-17-6P
RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of insecticidal and acaricidal 2-nitroguanidines)
IT 107-15-3, 1,2-Diaminoethane, reactions 110-60-1, 1,4-Diaminobutane
4245-76-5 7300-34-7 70258-18-3, 2-Chloro-5-chloromethylpyridine
105827-91-6, 2-Chloro-5-chloromethylthiazole

IT 220618-03-1P 220618-04-2P 220618-05-3P 220618-06-4P 220618-07-5P
220618-08-6P 220618-09-7P 220618-10-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. of insecticidal and acaricidal 2-nitroguanidines)

ANSWER 3 OF 33 CARBIUS COPYRIGHT 1999 ACS

ANSWER 3 OF 35 C
BN 1600-3030 GABRIUS

AN 1999:790
DN 130-E3438

DN 130-52438
TI Method for producing nitroguanidine derivatives

IN Maienfisch, Peter; Widmer, Hansjurg

IN Mainflrisch, 1991; Novartis-Erfindungen Verwaltungsgesellschaft
PA Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft

NOVEMBER

M.B.H.

so PCT Int. Appl., 23 pp.

CODEN: PIXXD2

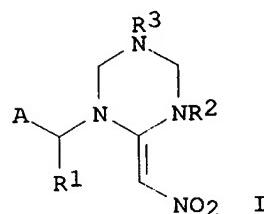
DT Patent

LA German

IC ICM C07D213-61

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9856764	A1	19981217	WO 1998-EP3358	19980605
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9884362	A1	19981230	AU 1998-84362	19980605
PRAI	CH 1997-1423		19970609		
	WO 1998-EP3358		19980605		
OS	MARPAT 130:52438				
	GI				



AB ACHR1NHC(NHR2):NNO2 [R1 = H, alkyl; R2 = H, alkyl, cycloalkyl, CH2B; A = (un)substituted arom. or non-arom. monocyclic or bicyclic or heterocyclic;

B = Ph, 3-pyridyl or thiazolyl which are optionally substituted by one to three substituents] were prep'd. by hydrolyzing a triazine I [R3 = (un)substituted alkyl, cycloalkyl, Ph, CH2Ph] at pH 7-14. The products are suitable as intermediates for the prodn. of pesticide mixts. Thus, I [A = 2-chloro-5-pyridyl, R1 = H, R2 = Me, R3 = Pr] was treated with

NaHCO3

in aq. MeOH to give ACH2NHC(NHMe):NNO2.

ST nitroguanidine prep'n pesticide intermediate; nitroiminotriazine hydrolysis

IT Pesticides

(prep'n. of nitroguanidine derivs. from nitroiminotriazines as intermediates for pesticides)

IT 136516-19-3 141856-78-2 141856-79-3

RL: RCT (Reactant)

(prep'n. of nitroguanidine derivs. from nitroiminotriazines as intermediates for pesticides)

IT 131748-47-5P 131748-50-0P 131748-56-6P 131748-59-9P 135018-04-1P
 135018-15-4P 135018-16-5P 135018-17-6P 135018-18-7P 135018-19-8P
 135018-20-1P 140920-89-4P

RL: SPN (Synthetic preparation); PREP (Preparation)

(prep'n. of nitroguanidine derivs. from nitroiminotriazines as intermediates for pesticides)

L7 ANSWER 4 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1998:672535 CAPLUS

DN 129:275911

TI Preparation of heterocycl-substituted nitroguandines.

IN Wollweber, Detlef; Kramer, Wolfgang; Rivadeneira, Eric

PA Bayer Aktiengesellschaft, Germany

SO PCT Int. Appl., 37 pp.

CODEN: PIXXD2

DT Patent

LA German

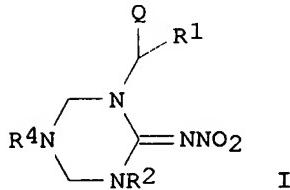
IC ICM C07D277-32

ICS C07D213-61; C07D405-06; C07D417-14

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9842690	A1	19981001	WO 1998-EP1456	19980313
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	DE 19712411	A1	19981001	DE 1997-19712411	19970325
	AU 9872068	A1	19981020	AU 1998-72068	19980313
PRAI	DE 1997-19712411	19970325			
	WO 1998-EP1456	19980313			
OS	CASREACT 129:275911; MARPAT 129:275911				
GI					



AB QR1CHNHC(:NNO₂)NHR₂ [R₁ = H, alkyl; R₂ = H, alkyl, cycloalkyl, CH₂R₃; R₃ =

alkenyl, alkynyl, Ph, cyanophenyl, nitrophenyl, halophenyl, pyridyl, thiazolyl, etc.; Q = (substituted) (arom.) mono- or bicyclic heterocyclyl], were prep'd. by treatment of triazacyclohexanes [I; R₄ = (substituted) alkyl, cycloalkyl, Ph, PhCH₂, heterocyclylmethyl; other variables as above] with urea in the presence of solvent. Thus, 1-(2-chlorothiazol-5-ylmethyl)-2-nitroimino-3,5-dimethyl-1,3,5-triazacyclohexane was refluxed 8 h with urea in isobutanol to give 84% 1-(2-chlorothiazol-5-ylmethyl)-2-nitro-3-methylguanidine. Guanidine heterocyclyl nitro prepns; chlorothiazolylmethylnitromethylguanidine prepns; heterocyclylnitroguanidine prepns; guanidine heterocyclyl nitro prepns; chlorothiazolylmethylnitromethylguanidine prepns

ST 131748-59-9P
IT RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(prepn. of heterocyclyl-substituted nitroguandines)

IT 617-89-0, Furfurylamine 4245-76-5 136516-19-3 213967-55-6

RL: RCT (Reactant)

(prepn. of heterocyclyl-substituted nitroguandines)

IT 213967-54-5P

RL: SPN (Synthetic preparation); PREP (Preparation)

(prepn. of heterocyclyl-substituted nitroguandines)

L7 ANSWER 5 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1998:668115 CAPLUS

DN 129:290052

TI Process for the preparation of nitroguanidine derivatives starting from 2-nitroiminohexahydro-1,3,5-triazines in the presence of ammonia, primary

or secondary amines
IN Ebihara, Koichi; Ura, Daisuke; Miyamoto, Michihiko; Kaiho, Tatsuo
PA Mitsui Chemicals, Inc., Japan
SO Eur. Pat. Appl., 13 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-61

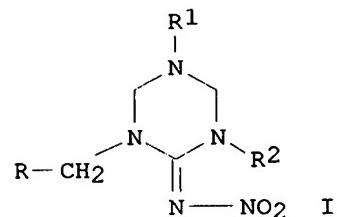
ICS C07D277-32; C07D307-14

ICA C07D401-06; C07D417-06; C07D407-06

CC 27-6 (Heterocyclic Compounds (One Hetero Atom))

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 869120	A1	19981007	EP 1998-105850	19980331
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	CN 1197064	A	19981028	CN 1998-108267	19980331
	JP 11236381	A2	19990831	JP 1998-86842	19980331
PRAI	JP 1997-80178		19970331		
	JP 1997-82838		19970401		
	JP 1997-223813		19970820		
	JP 1997-258968		19970924		
	JP 1997-347934		19971217		
OS	MARPAT 129:290052				
GI					



AB Described is a process, as a substitute for hydrolysis, for prepg. a nitroguanidine deriv., RCH₂NHC(:NNO₂)NHR₂ (R = 2-chloro-5-pyridyl, 2-chloro-5-thiazolyl, 2-, 3-tetrahydrofuryl, 2-methyl-4-tetrahydrofuryl; R₂ = Me, allyl), which comprises treating a triazine, I (R₁ = Me, benzyl, i-Pr, Et, t-Bu, cyclohexyl; R, R₂ = same as above), with NH₃, a primary amine or a secondary amine, or a salt thereof.

ST guanidine nitro prepn; iminohexahydrotriazine amination; triazine hydro aminative ring cleavage

IT 136516-18-2 165253-14-5 168688-97-9 195986-55-1 213967-55-6
214149-49-2 214149-50-5

RL: RCT (Reactant)

(aminative ring cleavage of)

IT 131748-47-5P 131748-59-9P 165252-70-0P 165253-05-4P 214149-48-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

L7 ANSWER 6 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1998:527323 CAPLUS

DN 129:136162

TI Preparation of thiazole derivatives.

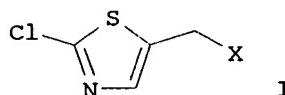
IN Piterna, Thomas; Maienfisch, Peter; Wadsworth, David John; Gsell, Laurenz; Rapold, Thomas; Szczepanski, Henry

PA Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft m.b.H.

SO PCT Int. Appl., 29 pp.

CODEN: PIXXD2
 DT Patent
 LA English
 IC ICM C07D277-32
 CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9832747	A1	19980730	WO 1998-EP297	19980120
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG	
	AU 9862929	A1	19980818	AU 1998-62929	19980120
PRAI	CH 1997-134		19970122		
	WO 1998-EP297		19980120		
OS	CASREACT 129:136162; MARPAT 129:136162				
GI					

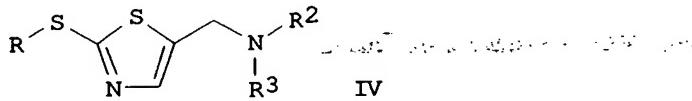
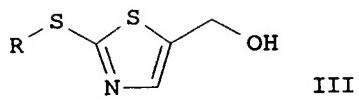
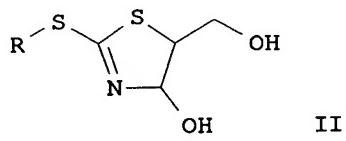
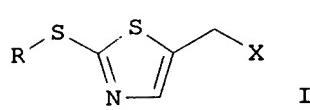


AB Title compds. (I; X = leaving group) were prepd. by (a) reacting 2-chloro-5-hydroxymethylthiazole with a sulfonylating agent to give I [X = OSO2A; A = alkyl, hydroxyalkyl, alkenyl, alkynyl, (substituted) aryl, etc.], or (b) reacting 2-chloro-5-chloromethylthiazole with an iodinating agent, preferably NaI to give I (X = iodo), or (c) reacting 2-chloro-5-methylthiazole with a brominating agent to give I (X = Br), etc. Thus, 2-chloro-5-methylthiazole was refluxed with dibenzoyl peroxide and NBS in CCl4 to give 5-bromomethyl-2-chlorothiazole.
 ST thiazole deriv prep; halomethylchlorothiazole prep; chlorothiazole halomethyl prep
 IT 210576-58-2P 210576-61-7P 210576-65-1P
 RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prep. of thiazole derivs.)
 IT 7143-01-3, Methanesulfonic anhydride 33342-65-3, 2-Chloro-5-methylthiazole 153719-38-1 192439-48-8
 RL: RCT (Reactant)
 (prep. of thiazole derivs.)
 IT 105827-91-6P, 2-Chloro-5-chloromethylthiazole 145015-15-2P,
 2-Chloro-5-hydroxymethylthiazole
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prep. of thiazole derivs.)
 IT 105829-23-0P 105843-36-5P 111988-48-8P 153719-23-4P 153719-32-5P
 153719-33-6P 171103-04-1P 209548-61-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prep. of thiazole derivs.)

L7 ANSWER 7 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1998:424238 CAPLUS
 DN 129:81723
 TI Preparation of thiazoles
 IN Pittnera, Thomas; Szczepanski, Henry; Maienfisch, Peter; Huter, Ottmar

Franz; Rapold, Thomas; Senn, Marcel; Gobel, Thomas; O'Sullivan, Anthony
 Cornelius
 PA Novartis A.-G., Switz.; Pitterna, Thomas; Szczepanski, Henry; Maienfisch,
 Peter; Huter, Ottmar Franz; Rapold, Thomas; Senn, Marcel; Gobel, Thomas;
 O'Sullivan, Anthony Cornelius
 SO PCT Int. Appl., 40 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 IC ICM C07D277-32
 ICS C07D277-16
 CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9827075	A1	19980625	WO 1997-EP7088	19971217
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9862056	A1	19980715	AU 1998-62056	19971217
	EP 946532	A1	19991006	EP 1997-954817	19971217
	R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, SE, IE, FI				
PRAI	CH 1996-3125		19961219		
	WO 1997-EP7088		19971217		
OS	MARPAT 129:81723				
GI					



AB The title compds. [I; R = (un)substituted C1-12 alkyl, C2-4 alkenyl, C2-4
 alkynyl, etc.; X = a leaving group] were prep'd. by a) reacting the
thiazoline II with a water-removing reagent; or b) reacting alc.
 III with a halogenating or a sulfonylating agent to form I [X = halo,
 sulfonate]; or c) reacting amine IV [R2, R3 = H, C1-6 alkyl, C3-6
 cycloalkyl, Ph, PhCH2] with haloc(O)OC1-8alkyl, haloC(O)Oaryl or
 haloC(O)OCH2Ph to obtain I [X = halo].

ST **thiazole** prep'n

IT 209473-20-1P 209473-21-2P

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prep'n. of **thiazoles**)

IT 192439-48-8P

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP

(Preparation)

(prep. of thiazoles)

IT 100-39-0, Benzyl bromide 765-34-4, Oxiranecarboxaldehyde 54895-19-1
 90197-29-8 209473-22-3
 RL: RCT (Reactant)
 (prep. of thiazoles)

L7 ANSWER 8 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1998:424237 CAPLUS

DN 129:95484

TI Preparation of thiazoles

IN Pitterna, Thomas; Szczeplanski, Henry; Maienfisch, Peter; Huter, Ottmar Franz; Rapold, Thomas; Senn, Marcel; Gobel, Thomas; O'Sullivan, Anthony Cornelius; Seifert, Gottfried

PA Novartis A.-G., Switz.

SO PCT Int. Appl., 47 pp.

CODEN: PIXXD2

DT Patent

LA English

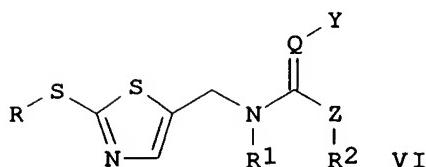
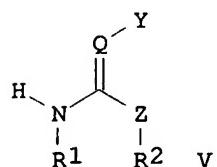
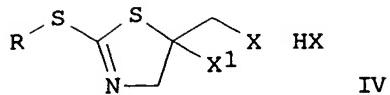
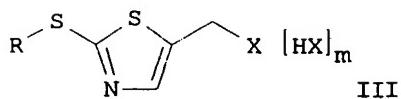
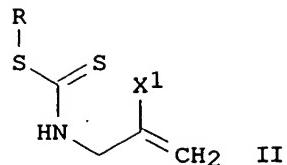
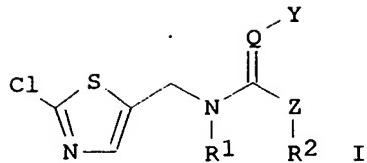
IC ICM C07D277-32

ICS C07D277-16; C07D417-06; C07D277-36

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9827074	A1	19980625	WO 1997-EP7087	19971217
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9857592	A1	19980715	AU 1998-57592	19971217
	EP 946531	A1	19991006	EP 1997-953841	19971217
	R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, SE, IE, FI				
	ZA 9711358	A	19980708	ZA 1997-11358	19971218
PRAI	CH 1996-3124		19961219		
	WO 1997-EP7087		19971217		
OS	CASREACT 129:95484; MARPAT 129:95484				
GI					



AB The title compds. [I; Q = CH, N; Y = NO₂, CN; Z = CHR₃, O, NR₃, S; R₁, R₂ = H, (un)substituted C₁-6 alkyl; R₁R₂ = alkylene which may addnl. contain a hetero atom selected from the group consisting of NR₅, O and S; R₃ = H, (un)substituted C₁-12 alkyl] were prep'd. by a) reacting dithiocarbamate II
 II [R = (un)substituted C₁-12 alkyl, C₂-4 alkenyl, C₂-4 alkynyl, etc.; X₁ = a leaving group] with a halogenating agent to form **thiazole** III [X = halo; m = 0-1], or by b) converting II by means of a halogenating agent into **thiazoline** IV, optionally c) converting IV into III, d) reacting III with the compd. V to form **thiazole** VI, e) or reacting IV with V to form **thiazole** VI, and f) treatment of compd. VI with chlorinating agent.

ST **thiazole** prep'n

IT	192439-34-2P	192439-36-4P	192439-37-5P	192439-38-6P	192439-39-7P
	192439-40-0P	192439-46-6P	192439-47-7P	192439-48-8P	192723-46-9P
	209548-64-1P	209548-65-2P	209548-66-3P	209548-67-4P	209548-68-5P
	209548-69-6P	209548-70-9P	209548-71-0P		

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of **thiazoles**)

IT	105829-23-0P	105843-36-5P	111988-48-8P	153719-23-4P	153719-32-5P
	153719-33-6P	171103-04-1P	209548-61-8P		

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of **thiazoles**)

IT	100-53-8, Benzyl mercaptan	14214-31-4, 2-Chloroallyl isothiocyanate			
	153719-38-1				

RL: RCT (Reactant)
 (prepn. of **thiazoles**)

L7 ANSWER 9 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1997:739397 CAPLUS

DN 128:44936

TI Systemic pesticide composition and method for treating citrus during pruning or grafting

IN Mizobe, Shinji; Miyata, Akiyoshi; Saito, Kenji

PA Nippon Soda Co., Ltd., Japan; Yamaguchi Prefecture
 SO Jpn. Kokai Tokkyo Koho, 7 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 IC ICM A01N033-18
 ICS A01G007-06; A01N037-20; A01N037-26; A01N037-34; A01N037-52;
 A01N043-40; A01N043-50; A01N043-68; A01N043-78; A01N047-40;
 A01N047-42; A01N047-44; C08L031-04
 CC 5-4 (Agrochemical Bioregulators)
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 09291002	A2	19971111	JP 1996-354437	19961219
PRAI	JP 1996-69290		19960229		
OS	MARPAT 128:44936				

AB A compn. for controlling pests on citrus fruits contains a systemic insecticide (I) of the formula RNAC(:YX)B at 0.1-10% by wt. and a vinyl acetate-type polymer; the compn. is applied to the cut surface in pruning or injected into grafts. In I, R = H, formyl, acetyl, C1-4 alkyl, 2-chloro-5-pyridylmethyl, or 2-chloro-5-thiazolylmethyl; A = H, C1-4 alkyl, or is bonded with B to form, e.g., an ethylene group; B =

C1-4 alkyl, SR₂ (R₂ = C1-4 alkyl), etc.; Y = N or CR₃ (R₃ = H or C1-4 alkyl); X = nitro or cyano. Such formulations may also contain a systemic antimicrobial agent. Thus, applying a compn. contg. acetamiprid 2, vinyl acetate adhesive 80, and surfactant, etc. 18% to the cut surface of a Citrus unshiu branch prevented damage by citrus leaf miner.

ST insecticide formulation pest control citrus; fungicide insecticide formulation citrus; chloronicotinyl insecticide formulation citrus

IT Pesticide formulations
 (contg. systemic insecticides for citrus treatment during pruning or grafting)

IT Phyllocnistis citrella
 (pesticide compns. and method for controlling leaf miner damage)

IT Binders
 Citrus
 Grafting (biological)
 Satsuma
 (pesticide compns. and method for treating citrus during pruning or grafting)

IT Fungicides
 (systemic fungicides; pesticide compns. and method for treating citrus during pruning or grafting)

IT Insecticides
 (systemic; pesticide compns. and method for treating citrus during pruning or grafting)

IT 9003-20-7, Vinyl acetate resin
 RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
 (pesticide compns. and method for treating citrus during pruning or grafting)

IT 23564-05-8, Topsin M 138261-41-3, Imidacloprid 160430-64-8,
 Acetamiprid
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BIOL (Biological study); USES (Uses)
 (pesticide compns. and method for treating citrus during pruning or grafting)

L7 ANSWER 10 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1997:429424 CAPLUS
 DN 127:46495
 TI Insect pest control method using liquid containing nonorganophosphorus insecticide
 IN Sekiyama, Atsuo

PA Takeda Seiyaku K. K., Japan
SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM A01N025-00

ICS A01N043-54; A01N043-62; A01N043-64

CC 5-4 (Agrochemical Bioregulators)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 09124402	A2	19970513	JP 1995-303586	19951027
OS	MARPAT 127:46495				
AB	Before transplanting vegetable or flowering plant seedlings, .apprx.0.1-20 g/m ² of a pesticidal liq.is applied to the nursery bed; the liq. contains 200-2000 ppm of a permeable insecticide R1R2NCR3:YX (I), where R1 = H, C1-6 alkyl, formyl, C1-4 alkylcarbonyl, C6-10 arylcarbonyl, C1-4 alkylsulfonyl, or a C1-3 alkyl substituted with a 3-6-membered heterocyclic group contg. N; R2 = H, hydrocarbon group, divalent group bonded with R3; R3 = hydrocarbon group, SR4 or NR5R6 (R4, R5, and R6 have the same meanings as R1) , or R3 is bonded with R2; Y = N, CH, hydrocarbon group; X = electron-withdrawing group. By applying a high rate of I, complete control can be achieved for .gtoreq.1 mo, and labor can be decreased markedly by giving seedlings water in which the insecticide is mixed during irrigation. Thus, applying 30 mL/seedling of a liq. contg. 333 ppm N-(2-chloro-5-thiazolylmethyl)-N'-methyl-N''- nitroguanidine, 1 day before planting, completely controlled cotton aphid for 41 days.				
ST	insecticide nonorganophosphorus pest control seedling transplanting; thiazolylmethyl nitroguanidine deriv cotton aphid control				
IT	Aphis gossypii Flower Insecticides Seedling Vegetable (pest control in beds for transplanting vegetable or flowering plant seedlings with liq. contg. nonorganophosphorus insecticide)				
IT	120738-89-8 131748-59-9 138261-41-3 160430-64-8 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BIOL (Biological study); USES (Uses) (pest control in beds for transplanting vegetable or flowering plant seedlings with liq. contg. nonorganophosphorus insecticide)				

=> d 17 11-20 all

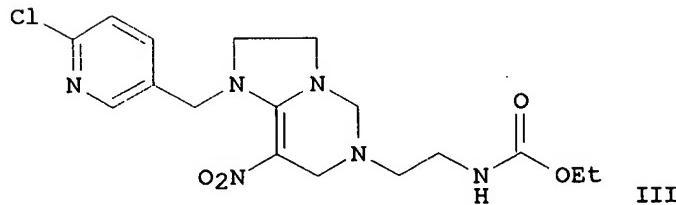
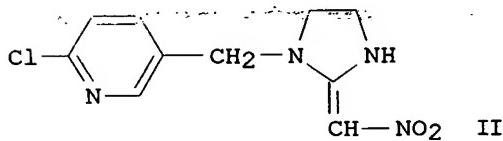
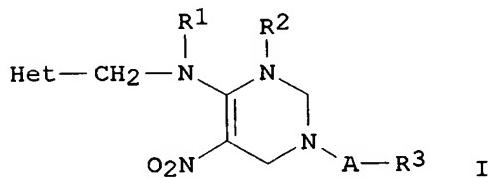
L7 ANSWER 11 OF 33 CAPLUS COPYRIGHT 1999 ACS
AN 1997:175073 CAPLUS
DN 126:171588
TI Preparation of heteroarylmethylisoureas and related compounds.
IN Uneme, Hideki; Konobe, Masato; Ishizuka, Hitoshi; Kamiya, Yasuo
PA Takeda Chemical Industries, Ltd., Japan; Uneme, Hideki; Konobe, Masato;
Ishizuka, Hitoshi; Kamiya, Yasuo
SO PCT Int. Appl., 48 pp.
CODEN: PIXXD2
DT Patent
LA English
IC ICM C07D277-28
ICS C07D213-38; C07D209-48; C07C273-18; C07C275-70; C07C277-08;
C07C279-36
CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9700867	A1	19970109	WO 1996-JP1694	19960619
	W: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CZ, EE, GE, HU, IL, IS, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, AM, AZ, BY, KG, KZ, MD				
	RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	CA 2220094	AA	19970109	CA 1996-2220094	19960619
	AU 9661374	A1	19970122	AU 1996-61374	19960619
	CN 1188475	A	19980722	CN 1996-194977	19960619
	EP 873325	A1	19981028	EP 1996-918850	19960619
	R: CH, DE, ES, FR, GB, IT, LI, PT				
	BR 9608892	A	19990615	BR 1996-8892	19960619
	JP 09067342	A2	19970311	JP 1996-162230	19960621
	JP 10120666	A2	19980512	JP 1996-304542	19961115
PRAI	JP 1995-158199		19950623		
	JP 1995-300278		19951117		
	WO 1996-JP1694		19960619		
	JP 1996-226595		19960828		
OS	CASREACT 126:171588; MARPAT 126:171588				
AB	Q(CH ₂) _n R ₂ C(OR ₁):NX [R ₁ = (substituted) hydrocarbyl; R ₂ = H, (substituted) hydrocarbyl; Q = (substituted) heterocyclyl; X = electron attracting group; n = 0, 1], were prep'd. by (A) reacting R ₁ O(N ₂ N)C:NX (I) with Q(CH ₂) _n NHR ₂ (II) or (B) reacting I with Y ₁ COACOY ₂ (Y ₁ , Y ₂ = leaving group) followed by treatment of the product with II. Thus, O-methyl-N-nitro-N'- phthaloylisourea (prepn. given) in MeOH was treated over 15 min. with 5-aminomethyl-2-chlorothiazole at 0.degree. and the mixt. was stirred 30 min. at room temp. to give 85% O-methyl-N-(2-chloro-5- thiazolylmethyl)-N'-nitroisourea. The latter in H ₂ O was treated dropwise with aq. MeNH ₂ followed by stirring for 14 h at room temp. to give 92% 1-(2-chloro-5-thiazolylmethyl)-3-methyl-2- nitroguanidine.				
ST	thiazolylmethylnitrosoisourea prep'n; heteroaryl methylisourea prep'n; pyridylmethylnitroguanidine prep'n				
IT	187149-02-6P	187149-03-7P			
	RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. of heteroaryl methylisoureas and related compds.)				
IT	131748-47-5P	131748-59-9P			
	RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation) (prepn. of heteroaryl methylisoureas and related compds.)				
IT	74-89-5, Methylamine, reactions 88-95-9, Phthaloyl chloride 24285-39-0, O-Methylisourea sulfate 97004-04-1, 5-Aminomethyl-2- chloropyridine 120740-08-1, 5-Aminomethyl-2-chlorothiazole RL: RCT (Reactant) (prepn. of heteroaryl methylisoureas and related compds.)				
IT	57538-27-9P	187149-01-5P			
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. of heteroaryl methylisoureas and related compds.)				
L7	ANSWER 12 OF 33 CAPLUS COPYRIGHT 1999 ACS				
AN	1995:957959 CAPLUS				
DN	124:8836				
TI	Substituted 1,2,3,4-tetrahydro-5-nitropyrimidines and their preparation and use as pesticides				
IN	Krueger, Bernd-Wieland; Uhr, Hermann; Kanellakopulos, Johannes; Gesing, Ernst R. F.; Wolf, Hilmar; Turberg, Andreas; Mencke, Norbert; Erdelen, Christoph; Wachendorff-Neumann, Ulrike; Hartwig, Juergen				
PA	Bayer A.-G., Germany				

SO Ger. Offen., 28 pp.
 CODEN: GWXXBX
 DT Patent
 LA German
 IC ICM C07D417-12
 ICS C07D401-12; C07D487-04; C07D498-06; A01N043-90; A01N043-54;
 A01N047-12; A01N047-18; A01N047-10; A01N047-28
 ICI C07D417-12, C07D239-06, C07D277-32; C07D401-12, C07D213-61, C07D239-06;
 C07D487-04, C07D239-00, C07D233-00; C07D487-04
 CC 28-16 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4401635	A1	19950727	DE 1994-4401635	19940121
	WO 9519977	A1	19950727	WO 1995-EP58	19950109
	W: AU, BB, BG, BR, BY, CA, CN, CZ, FI, HU, JP, KR, KZ, LK, MX, NO, NZ, PL, RO, RU, SK, UA, US RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CE, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9514161	A1	19950808	AU 1995-14161	19950109
	AU 697889	B2	19981022		
	EP 740666	A1	19961106	EP 1995-905616	19950109
	R: BE, CH, DE, ES, FR, GB, IT, LI, NL				
	CN 1138860	A	19961225	CN 1995-191285	19950109
	JP 09507670	T2	19970805	JP 1995-519313	19950109
	BR 9506532	A	19970916	BR 1995-6532	19950109
	ZA 9500462	A	19950928	ZA 1995-462	19950120
	US 5869491	A	19990209	US 1996-676272	19960712
PRAI	DE 1994-4401635	19940121			
	WO 1995-EP58	19950109			
OS	CASREACT 124:8836; MARPAT 124:8836				
GI					



AB Title compds. are claimed, specifically I [Het = (un)substituted pyridyl or thiazolyl; R1, R2 = Cl-4 alkyl; or R1R2 form (un)substituted

satd. 5- or 6-membered ring with optional O or N atoms; A = (un)substituted alkylene or cycloalkylene, the former optionally interrupted by O, S, alkylimino, or arylimino; R3 = OC(:X)R6, NR7C(:X)R6, SC(:X)R6; R6 = (un)substituted alkyl, cycloalkyl, alkenyl, aryl, (di)(alkyl)amino, etc.; X = O, S; R7 = H, C1-4 alkyl]. The compds. are useful as pesticides and ectoparasiticides, and esp. as insecticides.

For

example, cyclocondensation of (chloropyridinylmethyl)(nitromethylene)imida zolidine II with H₂NCH₂CH₂NHCO₂Et and formaldehyde in refluxing EtOH gave 82% title compd. III. In an in vitro test against Ctenocephalides felis, III showed 100% activity at 10 ppm. Addnl. biol. and synthetic examples are given.

ST tetrahydronitropyrimidine prepn pesticide; pyrimidine tetrahydronitro prepn ectoparasiticide; imidazolidinopyrimidine prepn insecticide acaricide nematocide

IT Acaricides
Insecticides
Nematocides
Pesticides
(prepns. of tetrahydronitropyrimidine derivs. as pesticides)

IT Parasiticides
(ecto-, prepns. of tetrahydronitropyrimidine derivs. as pesticides)

IT 113044-83-0P 170982-30-6P 170982-31-7P 170982-32-8P 170982-33-9P
170982-34-0P 170982-35-1P 170982-36-2P 170982-37-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(intermediate; prepns. of tetrahydronitropyrimidine derivs. as pesticides)

IT 170982-15-7P 170982-16-8P 170982-17-9P 170982-18-0P 170982-19-1P
170982-20-4P 170982-21-5P 170982-22-6P 170982-23-7P 170982-24-8P
170982-25-9P 170982-26-0P 170982-27-1P 170982-28-2P 170982-29-3P
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepns. of tetrahydronitropyrimidine derivs. as pesticides)

IT 50-00-0, Formaldehyde, reactions 78-96-6, 1-Amino-2-propanol
108-24-7,
Acetic anhydride 111-36-4, n-Butyl isocyanate 156-87-6,
3-Amino-1-propanol 36553-29-4 101336-63-4 120738-89-8
RL: RCT (Reactant)
(starting material; prepns. of tetrahydronitropyrimidine derivs. as pesticides)

L7 ANSWER 13 OF 33 CAPLUS COPYRIGHT 1999 ACS
AN 1994:191147 CAPLUS
DN 120:191147
TI Preparation of nitroguanidines as agrochemicals
IN Aoki, Isao; Minamida, Isao
PA Takeda Chemical Industries Ltd, Japan
SO Jpn. Kokai Tokkyo Koho, 24 pp.
CODEN: JKXXAF

DT Patent
LA Japanese
IC ICM C07C275-70
ICS C07C277-00; C07C279-34; C07C279-36; C07C335-40; C07D213-75;
C07D241-20; C07D265-06; C07D277-28; C07D277-32; C07D277-34;
C07D333-36; C07D417-12

ICA C07D279-06
CC 23-20 (Aliphatic Compounds)
Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05112521	A2	19930507	JP 1991-130805	19910322
OS	MARPAT	120:191147			

AB L-NH-A-G-C(NNO₂)-NR₂R₁ [I; R₁ = H, (un)substituted hydrocarbyl; R₂ = H, (un)substituted hydrocarbyl, (CH₂)_n-Y; Y = (un)substituted hydrocarbyl, substituted heterocyclyl; n = 1, 2; R₁R₂N = part of a ring; A = (un)substituted alkylene; G = O, S; L = electron-withdrawing group] are prep'd. A mixt. of 2-(nitroimino)-3-(p-tolylsulfonyl)tetrahydro-4H-1,3-thiazine and 2-(2-chlorophenyl)ethylamine in CHCl₃ was stirred at room temp. for 1 h to give I [L = p-tolylsulfonyl, A = (CH₂)₃, G = S, R₁ = H, R₂ = o-chlorophenethyl]. I were active as pesticides at 1-1000 ppm, preferably at 1-500 ppm. I may be used as herbicides, acaricides, insecticides, fungicides, etc. (no data).

ST nitroguanidine prepn agrochem; guanidine nitro prepn agrochem

IT Agrochemicals
(nitroguanidines)

IT	35089-65-7P	101250-97-9P	131748-47-5P	131748-48-6P	131748-49-7P
	131748-50-0P	131748-51-1P	131748-53-3P	131748-55-5P	131748-56-6P
	131748-58-8P	131748-59-9P	131748-60-2P	131748-61-3P	131748-65-7P
	131748-69-1P	131748-70-4P	131748-71-5P	131748-72-6P	131748-73-7P
	131748-74-8P	131748-75-9P	131748-76-0P	131748-77-1P	131748-78-2P
	131748-85-1P	131748-86-2P	131768-12-2P	131768-13-3P	135018-15-4P
	137589-63-0P	138149-89-0P	138149-90-3P	138149-92-5P	138149-95-8P
	141205-36-9P	141205-38-1P	141205-39-2P	141205-40-5P	141205-44-9P
	141205-46-1P	141205-49-4P	141205-50-7P	141205-51-8P	141205-52-9P
	141205-53-0P	141205-54-1P	141205-56-3P	141205-57-4P	141205-58-5P
	141205-60-9P	141205-61-0P	141205-63-2P	141205-64-3P	141205-65-4P
	141205-66-5P	141205-67-6P	141205-68-7P	141205-69-8P	141205-70-1P
	141205-71-2P	141205-72-3P	141205-73-4P	141205-74-5P	141205-76-7P
	141205-77-8P	141205-78-9P	141205-79-0P	141205-80-3P	141205-81-4P
	141205-82-5P	141205-83-6P	141205-84-7P	141227-31-8P	141227-32-9P
	141227-33-0P	141227-34-1P	150221-67-3P	150221-68-4P	150221-69-5P
	150221-70-8P	150221-71-9P			

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, as agrochem.)

IT 150221-72-0P,

2-(Nitroimino)-3-(p-tolylsulfonyl)tetrahydro-4H-1,3-thiazine

150221-73-1P	150221-76-4P	150221-77-5P	150221-78-6P	150221-79-7P
150221-80-0P	150221-82-2P	150221-84-4P	150221-85-5P	150221-86-6P
150221-89-9P				

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, as intermediate for agrochems.)

IT 98-59-9, p-Toluenesulfonyl chloride 107-10-8, Propylamine, reactions
124-63-0, Methanesulfonyl chloride 1848-68-6,

2-Amino-5,6-dihydro-4H-1,3-oxazine 1885-14-9, Phenyl chlorocarbonate 7073-36-1,
2-Chloro-4-nitrobenzoyl chloride 13078-80-3, 2-(2-Chlorophenyl)ethylamine 27757-85-3, 2-Aminomethylthiophene 30480-64-9
97004-04-1 105827-90-5, 2-Nitroiminothiazolidine 120740-08-1,
(2-Chloro-5-thiazolyl)methylamine 150221-75-3,
2-Nitroimino-3-(phenylsulfonyl)thiazolidine 150221-81-1
150221-83-3, 2-Imino-3-phenylsulfonylthiazolidine 150221-87-7,
2-Nitroiminotetrahydro-4H-1,3-thiazine 150221-88-8, 2-Nitroiminotetrahydro-4H-1,3-oxazine

RL: RCT (Reactant)
(reaction of, in prepn. of agrochems.)

L7 ANSWER 14 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1994:127812 CAPLUS

DN 120:127812

TI Termite-proofing agent

IN Matsuda, Michihiko; Hatano, Renpei; Yano, Makio

PA Nippon Soda Co., Ltd., Japan

SO PCT Int. Appl., 11 pp.

CODEN: PIXXD2

DT Patent

LA Japanese

IC ICM A01N047-40

ICS A01N051-00

CC 5-4 (Agrochemical Bioregulators)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9325080	A1	19931223	WO 1993-JP755	19930604
	W: AU, BR, US			RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE	
	AU 9343545	A1	19940104	AU 1993-43545	19930604
	AU 670441	B2	19960718		
	JP 06056612	A2	19940301	JP 1993-158177	19930604
	EP 651945	A1	19950510	EP 1993-913481	19930604
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, MC, NL, PT, SE				
	BR 9306540	A	19980915	BR 1993-6540	19930604
PRAI	JP 1992-177395		19920611		
	WO 1993-JP755		19930604		
OS	MARPAT 120:127812				
AB	A termite-proofing agent contg. at least either a compd. represented by general formula I ($R_1CH_2NR_2CR_3:NR_4$) or a salt thereof as the active ingredient, wherein R_1 represents pyridyl, pyrazinyl or thiazolyl each of which may be substituted; R_2 represents H, optionally substituted alkyl, cycloalkyl or alkoxy; R_3 represents H or optionally substituted alkyl; and R_4 represents cyano or NO_2 .				
ST	termite control amine deriv				
IT	Insecticides (amine derivs., for termite control)				
IT	Termite (control of, amine derivs. for)				
IT	Agrochemical formulations (insecticide-contg., for termite control)				
IT	135410-03-6	135410-19-4	135410-20-7	135410-21-8	135410-40-1
	135410-42-3	135410-43-4	135410-52-5	135410-60-5	135410-81-0
	135410-83-2	135410-92-3	136479-54-4		
	RL: BIOL (Biological study) (insecticide compns. contg., for termite control)				

L7 ANSWER 15 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1993:580670 CAPLUS

DN 119:180670

TI Preparation of guanidine containing heterocycles as insecticides

IN Tsuboi, Shinichi; Moriya, Koichi; Hattori, Yumi; Sone, Shinzaburo;
Shibuya, Katsuhiko

PA Nihon Bayer Agrochem K.K., Japan

SO Eur. Pat. Appl., 12 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-61

ICS C07D277-32; C07D261-10; A01N043-40; A01N043-78; A01N043-80

CC 27-16 (Heterocyclic Compounds (One Hetero Atom))

Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 547451	A1	19930623	EP 1992-120735	19921204
	EP 547451	B1	19970305		
	R: BE, CH, DE, ES, FR, GB, IT, LI, NL				
	JP 05163242	A2	19930629	JP 1991-352861	19911217
	ES 2097856	T3	19970416	ES 1992-120735	19921204
	US 5304564	A	19940419	US 1992-987510	19921207
	ZA 9209716	A	19930614	ZA 1992-9716	19921215
	BR 9205027	A	19930622	BR 1992-5027	19921216
	CN 1073678	A	19930630	CN 1992-114469	19921217
	CN 1037737	B	19980318		
	HU 63306	A2	19930830	HU 1992-3996	19921217

US 5472971 A 19951205 US 1994-182957 19940118
 PRAI JP 1991-352861 19911217
 US 1992-987510 19921207
 OS MARPAT 119:180670
 AB Title compds. ZCHR1NH(CH₂)_nR2C(:NY)NR3R4 (I; Z = 2-chloro-5-pyridyl, 2-chloro-5-thiazolyl, 3-chloro-5-isoxazolyl; R₁ = H, C₁-4 alkyl; R₂ = H, C₁-4 alkyl, C₃-4 alkenyl, C₃-4 alkynyl, 2-chloro-5-pyridylmethyl; R₃, R₄ = H, halo, C₁-4 alkyl, C₃-4 alkenyl, C₃-4 alkynyl, (substituted) PhCH₂, etc.; Y = O₂N, NC; n = 2,3), are prepd. N-(6-chloro-3-pyridylmethyl)ethylenediamine, MeSC(:NNO₂)NH₂ and EtOH were stirred at 30.degree. until MeSH ceased to be generated to give I (Z = 6-chloro-3-pyridyl, R₁ = R₂ = R₃ = R₄ = H, Y = O₂N, n = 2) (II). In test on plant hoppers planted on rice plant, II at 200 ppm gave 100% control.
 ST heterocyclyl guanidine prepn insecticide
 IT Insecticides
 (guanidine contg. heterocycles)
 IT 150312-16-6P 150433-40-2P 150433-41-3P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of, as insecticide)
 IT 2986-25-6 101990-44-7
 RL: RCT (Reactant)
 (reaction of, in prepn. of insecticides)

 L7 ANSWER 16 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1993:533485 CAPLUS
 DN 119:133485
 TI Heterocyclic nematocides and acaricides.
 IN Matsuda, Michihiko; Takakusa, Nobuo; Yamamoto, Atsushi; Yano, Makio
 PA Nippon Soda Co, Japan
 SO Jpn. Kokai Tokkyo Koho, 6 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 IC ICM A01N047-40
 ICS A01N051-00
 CC 5-4 (Agrochemical Bioregulators)
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 05155722	A2	19930622	JP 1991-348450	19911205

OS MARPAT 119:133485
 AB Nematocides and acaricides contain R₁CH₂NR₂CR₃:NR₄ [I; R₁ = (un)substituted pyridyl, pyrazinyl or thiazolyl; R₂ = H, (un)substituted alkyl, cycloalkyl, alkoxy; R₃ = H, (un)substituted alkyl; R₄ = cyano, NO₂] as active ingredients. I (R₁ = 2-chloropyridin-5-yl, R₂ = R₃ = Me, R₄ = cyano) 20, higher alc. sulfate ester 5, diatomaceous earth 70, and silica 5 parts were mixed to give a wettable powder, which showed higher nematocidal activity than cartap.
 ST nematocide acaricide pyridine pyrazine thiazole
 IT Acaricides
 Nematocides
 (pyridines and pyrazines and thiazoles)
 IT 135410-20-7 135410-42-3 135410-81-0 135410-92-3 136479-54-4
 RL: BIOL (Biological study)
 (acaricide and nematocide)

 L7 ANSWER 17 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1993:141817 CAPLUS
 DN 118:141817
 TI Agrohorticultural insecticidal and bactericidal compositions containing amines.
 IN Matsuda, Michihiko; Takakusa, Nobuo; Yamamoto, Atsushi; Iwasa, Takao;

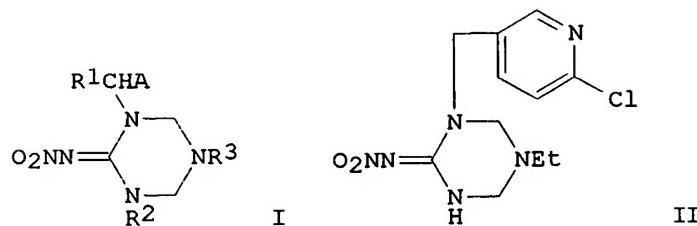
PA Hashimoto, Sho
 Nippon Soda Co., Ltd., Japan
 SO PCT Int. Appl., 19 pp.
 CODEN: PIXXD2
 DT Patent
 LA Japanese
 IC ICM A01N047-40
 ICS A01N051-00
 CC 5-2 (Agrochemical Bioregulators)
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9221241	A1	19921210	WO 1992-JP714	19920602
	W: BR, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE				
	JP 04360804	A2	19921214	JP 1991-159690	19910604
	JP 04368304	A2	19921221	JP 1991-167426	19910613
PRAI	JP 1991-159690		19910604		
	JP 1991-167426		19910613		
OS	MARPAT 118:141817				
AB	An amine deriv. (R1X)NR2CR3:NR4 (I) [R1 = (un)substituted pyridyl, pyrazyl, thiazolyl; X = (un)substituted alkylene, alkylidene; R2 = H, (un)substituted alkyl, alkenyl, YR5, etc.; Y = O, CO; R3 = H, (un)substituted alkyl, cycloalkyl; R4 = cyano, nitro; R5 = (un)substituted alkyl, aryl] is an insecticide and bactericide in combination with an ergosterol biosynthesis inhibitor, like triflumizole. A I [R1 = ergosterol biosynthesis inhibitor, like triflumizole. A I [R1 = 3-chloro-3-pyridiyl, X = CH ₂ , R2 = Me, R3 = Me, R4 = CN] mixt. with triflumizole showed potent insecticidal activities.				
ST	insecticide bactericide triflumizole amine deriv				
IT	Bactericides, Disinfectants, and Antiseptics				
	Insecticides				
	(amine deriv. mixts. with ergosterol biosynthesis inhibitors)				
IT	146543-66-0	146543-67-1	146543-68-2	146543-69-3	146543-70-6
	146543-71-7	146543-72-8	146543-73-9	146543-74-0	146543-75-1
	146543-76-2	146543-77-3	146543-78-4	146543-79-5	146543-80-8
	146543-81-9	146570-99-2	146571-01-9	146571-02-0	
RL	BIOL (Biological study)				
	(bactericide and insecticide)				

L7 ANSWER 18 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1992:469886 CAPLUS
 DN 117:69886
 TI Preparation of 2-(nitroimino)-1,3,5-triazacyclohexane pesticides
 IN Maienfisch, Peter; Kristiansen, Odd; Gsell, Laurenz
 PA Ciba-Geigy A.-G., Switz.
 SO Eur. Pat. Appl., 102 pp.
 CODEN: EPXXDW
 DT Patent
 LA German
 IC ICM C07D401-06
 ICS C07D417-06; A01N051-00
 ICI C07D401-06, C07D251-00, C07D213-00; C07D417-06, C07D277-00, C07D251-00
 CC 28-19 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 483055	A1	19920429	EP 1991-810757	19910926
	R: BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL				
	CN 1060468	A	19920422	CN 1991-109409	19910930
	CA 2052731	AA	19920406	CA 1991-2052731	19911003
	JP 04273863	A2	19920930	JP 1991-283521	19911003
	BR 9104300	A	19920602	BR 1991-4300	19911004

PRAI CH 1990-3219 19901005
 CH 1991-1648 19910604
 OS MARPAT 117:69886
 GI



AB Title compds. [I; R1 = H, alkyl; R2 = H, (cyclo)alkyl, CH2B; R3 = H, (substituted) (cyclo)alkyl alkenyl, alkynyl, Ph, PhCH2; A = (substituted) (arom.) (bicyclic) heterocyclyl; B = (substituted) Ph, pyridyl, thiazolyl], were prepd. Thus, 1-(2-chloropyrid-5-ylmethyl)-2-nitroguanidine, aq. H₂CO, aq. EtNH₂, and EtOH were heated at 50.degree. for 4 h to give title compd. II. II at 400 ppm gave >80% control of

Myzus persicae.

ST nitroiminotriazacyclohexane prepn pesticide; insecticide nitroiminotriazacyclohexane; acaricide nitroiminotriazacyclohexane; triazacyclohexane nitroimino prepн pesticide

IT Acaricides

Insecticides

Pesticides

(nitroimino)triazacyclohexanes)

IT	13423-25-4P	136516-17-1P	136516-18-2P	141856-77-1P	141856-78-2P
	141856-79-3P	141856-80-6P	141856-81-7P	141856-82-8P	141856-83-9P
	141856-84-0P	141856-85-1P	141856-86-2P	141856-87-3P	141856-88-4P
	141856-89-5P	141856-91-9P	141856-92-0P	141856-93-1P	141856-94-2P
	141856-95-3P	141856-96-4P	141856-97-5P	141856-98-6P	141856-99-7P
	141857-00-3P	141857-01-4P	141857-02-5P	141857-03-6P	141857-04-7P
	141857-05-8P	141857-06-9P	141857-07-0P	141857-08-1P	141857-09-2P
	141857-10-5P	141857-11-6P	141857-12-7P	141857-13-8P	141857-14-9P
	141857-15-0P	141857-16-1P	141857-17-2P	141857-18-3P	141857-19-4P
	141857-21-8P	141857-22-9P	141857-23-0P	142488-63-9P	142488-64-0P
	142488-65-1P	142488-66-2P	142488-67-3P		

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of, as pesticide)

IT	133258-70-5P	136516-16-0P	141856-46-4P	141856-47-5P	141856-48-6P
	141856-49-7P	141856-50-0P	141856-51-1P	141856-52-2P	141856-53-3P
	141856-54-4P	141856-55-5P	141856-56-6P	141856-57-7P	141856-59-9P
	141856-60-2P	141856-62-4P	141856-64-6P	141856-65-7P	141856-66-8P
	141856-67-9P	141856-68-0P	141856-69-1P	141856-70-4P	141856-71-5P
	141856-72-6P	141856-73-7P	141856-74-8P	141856-75-9P	141856-76-0P
	142488-68-4P	142488-69-5P			

RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of, as pesticide intermediate)

IT 50-00-0, Formaldehyde, reactions 74-89-5, Methylamine, reactions 107-10-8, Propylamine, reactions 556-88-7, 2-Nitroguanidine 4245-76-5,

1-Methyl-2-nitroguanidine

RL: RCT (Reactant)

(reaction of, in prepn. of (nitroamino)triazacyclohexane pesticide)

IT 75-04-7, Ethylamine, reactions 107-08-4, Propyl iodide 70258-18-3,

2-Chloro-5-chloromethylpyridine 131748-56-6
RL: RCT (Reactant)
(reaction of, in prepn. of pesticide)

L7 ANSWER 19 OF 33 CAPLUS COPYRIGHT 1999 ACS
AN 1992:448540 CAPLUS

DN 117:48540

TI Preparation of N-(thiazolylalkyl)-N'-cyanoguanidine or
-S-methylisothiourea derivatives and analogs as insecticides
IN Ishimitsu, Keiichi; Kishimoto, Takashi; Oishi, Haruhito; Yamada, Tomio;
Hatano, Renpei; Takakusa, Nobuo
PA Nippon Soda Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 11 pp.
CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C07D277-28

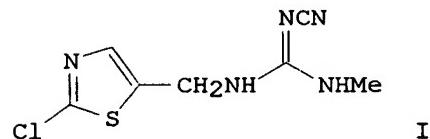
ICS A01N047-42; A01N047-44; C07D277-32; C07D277-34; C07D277-50

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 04029983	A2	19920131	JP 1990-132537	19900524
OS	MARPAT 117:48540				

GI



AB R1XNR2C(:NCN)R3 [R1 = (un)substituted 4- or 5-thiazolyl; X = (un)substituted alkylene, heteroatom; R2 = H, (un)substituted alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, YR4, NR5R6; Y = O, S(O)1, CO, CO2; l = 0-2; or XR2 forms a ring optionally contg. hetero atoms; R3 = SR7, NR8R9; R4-R8 = H, (un)substituted alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl; R9 = groups listed in R4-R8, ZR10, NR11R12; Z = groups listed in Y; R10 = groups listed in R4-R8; R11, R12 = groups listed in R4-R8, WR13; W = groups listed in Y; R13 = groups listed in R4-R8; or R8R9 forms a ring optionally contg. hetero atoms] are prepd. Thus, refluxing 3.0 g NCNC(SMe)2 and 2-chloro-5-thiazolylmethylamine in EtOH for 1 h with stirring and condensation of the resulting N-(2-chloro-5-thiazolylmethyl)-N'-cyano-S-methylisothiourea with MeNH2 in EtOH under reflux gave title compd. I. I at 125 ppm killed 100% cotton aphid on cucumber seedlings.

ST thiazolylalkylcyanoguanidine prepn insecticide; cyanoguanidine
thiazolyl alkyl insecticide

IT Insecticides

(N-(thiazolylalkyl)-N'-cyanoguanidine or -S-methylisothiourea
derivs. and analogs)

IT 131748-79-3P 141761-53-7P 141761-54-8P 141761-55-9P 141761-56-0P
RL: AGR (Agricultural use); BAC (Biological activity or effector, except
adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
(Preparation); USES (Uses)

(prepn. of, as insecticide)

IT 74-89-5, Methylamine, reactions 10191-60-3 120740-08-1
RL: RCT (Reactant)

(reaction of, in prepn. of insecticide)

L7 ANSWER 20 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1992:426353 CAPLUS
 DN 117:26353
 TI Process for the preparation of nitroguanidine derivatives
 IN Maienfisch, Peter; Kristiansen, Odd; Gsell, Laurenz
 PA Ciba-Geigy A.-G., Switz.
 SO Eur. Pat. Appl., 52 pp.
 CODEN: EPXXDW

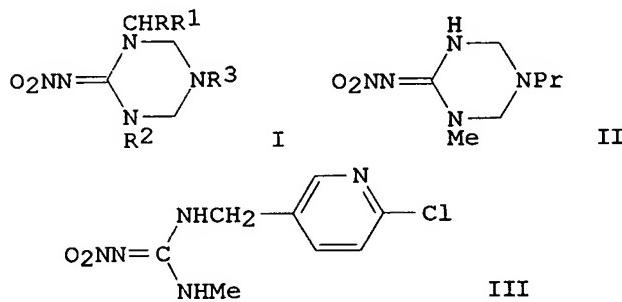
DT Patent
 LA German

IC ICM C07D213-61
 ICS A01N043-40; C07D213-40; C07D277-32; A01N043-78
 CC 27-16 (Heterocyclic Compounds (One Hetero Atom))

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 483062	A2	19920429	EP 1991-810795	19911015
	EP 483062	A3	19921028		
	EP 483062	B1	19960918		
	R: BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL ES 2091899 US 5245040 IL 99801 CA 2053954 JP 04330049	T3	19961116	ES 1991-810795 US 1991-777856 IL 1991-99801 CA 1991-2053954 JP 1991-303960	19911015 19911016 19911018 19911022 19911023
PRAI	CH 1990-3395		19901024		
OS	CASREACT 117:26353; MARPAT 117:26353				

GI



AB O₂NN:C(NHR₂)NHCHR₁ (R = heterocyclic; R₁ = H, alkyl; R₂ = H, alkyl, aralkyl, cycloalkyl) were prep'd. by hydrolyzing triazines I [R₃ = (un)substituted alkyl, cycloalkyl, Ph, CH₂Ph]. Thus, 17.1 g O₂NN:C(NH₂)NHMe was treated with PrNH₂ and aq. CH₂OH, to give 26.9 g triazine II. Treatment of 20.1 g II with 16.2 g 2-chloro-5-chloromethylpyridine to give 17.4 g I (R = 2-chloro-5-pyridyl, R₁, R₂ = H,

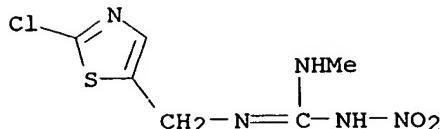
H, R₃ = Pr) which (4.25 g) was hydrolyzed with HCl-MeOH to give 2.51 g nitroguanidine III.

ST nitroguanidine pyridylmethyl thiazolylmethyl;
 pyridylmethylnitroguanidine; thiazolylmethylnitroguanidine

IT 141856-78-2P 141856-84-0P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. and hydrolysis of)

IT 141856-46-4P 141856-49-7P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. and reaction of, with chloro(chloromethyl)pyridine)

IT 35089-65-7P 131748-47-5P 131748-50-0P 131748-53-3P 131748-56-6P
 131748-59-9P 133258-61-4P 133258-65-8P 133258-66-9P 133258-70-5P
 134323-25-4P 135018-04-1P 135018-10-9P 135018-11-0P 135018-14-3P
 135018-15-4P 135018-16-5P 135018-17-6P 135018-18-7P 135018-19-8P
 135018-20-1P 136516-16-0P 136516-17-1P 136516-18-2P 140920-89-4P



L5 ANSWER 3 OF 4 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1993:59590 CAPLUS

DOCUMENT NUMBER: 118:59590

TITLE: Heterocyclic amidine derivatives
[(heteroaryl methyl)nitroguanidine derivatives] and
their use as pesticides (insecticides and acaricides)

INVENTOR(S): Kristiansen, Odd; Gsell, Laurenz; Maienfisch, Peter

PATENT ASSIGNEE(S): Ciba-Geigy A.-G., Switz.

SOURCE: Eur. Pat. Appl., 63 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 507736	A1	19921007	EP 1992-810225	19920326
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, PT				
US 5223520	A	19930629	US 1992-858910	19920327
CA 2064920	AA	19921005	CA 1992-2064920	19920402
JP 05117237	A2	19930514	JP 1992-109199	19920402
AU 9214036	A1	19921008	AU 1992-14036	19920403
CN 1065456	A	19921021	CN 1992-102347	19920403
HU 60721	A2	19921028	HU 1992-1140	19920403
BR 9201197	A	19921201	BR 1992-1197	19920403
ZA 9202450	A	19930405	ZA 1992-2450	19920403
PRIORITY APPLN. INFO.:		CH 1991-1004		19910404

OTHER SOURCE(S): CASREACT 118:59590; MARPAT 118:59590

AB Some heterocyclic amidine derivs. and a process for their prepn.
are claimed. The use of these compds. as pesticides (insecticides and
acaricides) is claimed. Condensation of 1-amino-2-[(2-chloro-5-
pyridyl)methyl]methylamino]-2-nitroethene with DMF di-Et acetal gave the
acyclic amidine I. I had activity against Nilaparvata lugens,

Nephrotettix

cincticeps, Bemisia tabaci and Ctenocephalides felis (flea; systemic).

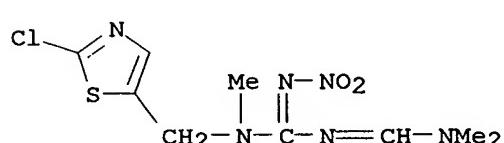
IT 145368-72-5P 145368-73-6P 145368-74-7P

145368-81-6P 145368-82-7P 145368-83-8P

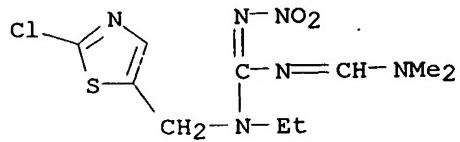
RL: AGR (Agricultural use); BAC (Biological activity or effector, except
adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
(Preparation); USES (Uses)
(prepn. of, as pesticide)

RN 145368-72-5 CAPLUS

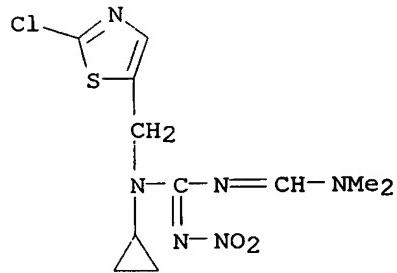
CN Guanidine,

N-[(2-chloro-5-thiazolyl)methyl]-N'-(dimethylamino)methylene)-
N-methyl-N''-nitro- (9CI) (CA INDEX NAME)

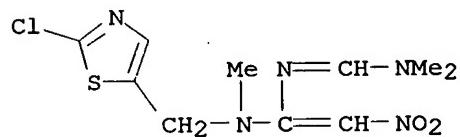
RN 145368-73-6 CAPLUS
CN Guanidine,
N-[(2-chloro-5-thiazolyl)methyl]-N'-(dimethylamino)methylene)-
N-ethyl-N''-nitro- (9CI) (CA INDEX NAME)



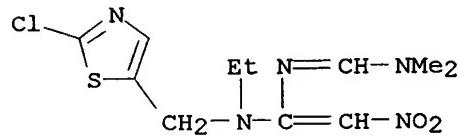
RN 145368-74-7 CAPLUS
CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N-cyclopropyl-N'-(dimethylamino)methylene)-N''-nitro- (9CI) (CA INDEX NAME)



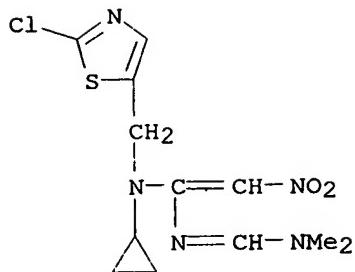
RN 145368-81-6 CAPLUS
CN Methanimidamide, N'-[1-[(2-chloro-5-thiazolyl)methyl]methylamino]-2-nitroethenyl]-N,N-dimethyl- (9CI) (CA INDEX NAME)



RN 145368-82-7 CAPLUS
CN Methanimidamide, N'-[1-[(2-chloro-5-thiazolyl)methyl]ethylamino]-2-nitroethenyl]-N,N-dimethyl- (9CI) (CA INDEX NAME)



RN 145368-83-8 CAPLUS
CN Methanimidamide,
N'-[1-[(2-chloro-5-thiazolyl)methyl]cyclopropylamino]-2-nitroethenyl]-N,N-dimethyl- (9CI) (CA INDEX NAME)



L5 ANSWER 4 OF 4 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1992:426353 CAPLUS

DOCUMENT NUMBER: 117:26353

TITLE: Process for the preparation of nitroguanidine derivatives

INVENTOR(S): Maienfisch, Peter; Kristiansen, Odd; Gsell, Laurenz

PATENT ASSIGNEE(S): Ciba-Geigy A.-G., Switz.

SOURCE: Eur. Pat. Appl., 52 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 483062	A2	19920429	EP 1991-810795	19911015
EP 483062	A3	19921028		
EP 483062	B1	19960918		
	R: BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL ES 2091899	19961116	ES 1991-810795	19911015
US 5245040	A	19930914	US 1991-777856	19911016
IL 99801	A1	19960723	IL 1991-99801	19911018
CA 2053954	AA	19920425	CA 1991-2053954	19911022
JP 04330049	A2	19921118	JP 1991-303960	19911023
			CH 1990-3395	19901024

PRIORITY APPLN. INFO.:

OTHER SOURCE(S): CASREACT 117:26353; MARPAT 117:26353

AB O₂NN:C(NHR₂)NHCHRR₁ (R = heterocyclic; R₁ = H, alkyl; R₂ = H, alkyl, aralkyl, cycloalkyl) were prep'd. by hydrolyzing triazines I [R₃ = (un)substituted alkyl, cycloalkyl, Ph, CH₂Ph]. Thus, 17.1 g O₂NN:C(NH₂)NHMe was treated with PrNH₂ and aq. CH₂OH to give 26.9 g triazine II. Treatment of 20.1 g II with 16.2 g 2-chloro-5-chloromethylpyridine to give 17.4 g I (R = 2-chloro-5-pyridyl, R₁, R₂ = H,

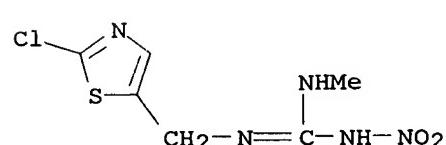
H, R₃ = Pr) which (4.25 g) was hydrolyzed with HCl-MeOH to give 2.51 g nitroguanidine III.

IT 131748-59-9P 135018-15-4P 135018-16-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

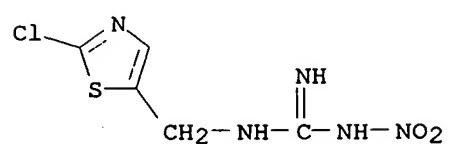
RN 131748-59-9 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitro- (9CI)
(CA INDEX NAME)

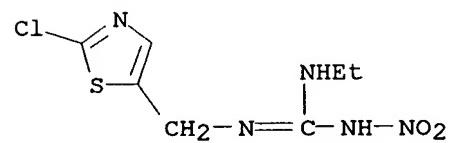


RN 135018-15-4 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-nitro- (9CI) (CA INDEX
NAME)



RN 135018-16-5 CAPLUS
CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-ethyl-N''-nitro- (9CI)
(CA INDEX NAME)



141856-47-5P	141856-48-6P	141856-50-0P	141856-51-1P	141856-52-2P
141856-53-3P	141856-54-4P	141856-55-5P	141856-56-6P	141856-57-7P
141856-58-8P	141856-59-9P	141856-60-2P	141856-61-3P	141856-62-4P
141856-63-5P	141856-64-6P	141856-65-7P	141856-66-8P	141856-67-9P
141856-68-0P	141856-69-1P	141856-70-4P	141856-71-5P	141856-72-6P
141856-73-7P	141856-74-8P	141856-75-9P	141856-76-0P	141856-77-1P
141856-79-3P	141856-80-6P	141856-81-7P	141856-82-8P	141856-83-9P
141856-85-1P	141856-86-2P	141856-87-3P	141856-88-4P	141856-89-5P
141856-90-8P	141856-91-9P	141856-92-0P	141856-93-1P	141856-94-2P
141856-95-3P	141856-96-4P	141856-97-5P	141856-98-6P	141856-99-7P
141857-00-3P	141857-01-4P	141857-02-5P	141857-03-6P	141857-04-7P
141857-05-8P	141857-06-9P	141857-07-0P	141857-08-1P	141857-09-2P
141857-10-5P	141857-11-6P	141857-12-7P	141857-13-8P	141857-14-9P
141857-15-0P	141857-16-1P	141857-17-2P	141857-18-3P	141857-19-4P
141857-20-7P	141857-21-8P	141857-22-9P	141857-23-0P	141857-24-1P
141857-25-2P	141857-26-3P	141857-27-4P	141857-28-5P	

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

IT 556-88-7, Nitroguanidine 4245-76-5

RL: RCT (Reactant)

(reaction of, with amines and formaldehyde)

IT 62-53-3, Aniline, reactions 74-89-5, Methylamine, reactions 107-10-8,
Propylamine, reactions

RL: RCT (Reactant)

(reaction of, with nitroguanidine and formaldehyde)

IT 50-00-0, Formaldehyde, reactions

RL: RCT (Reactant)

(reaction of, with nitroguanidines and amines)

IT 70258-18-3, 2-Chloro-5-chloromethylpyridine

RL: RCT (Reactant)

(reaction of, with nitroiminotriazacyclohexanes)

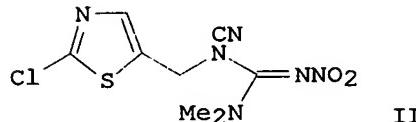
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L7 ANSWER 21 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1992:214489 CAPLUS
 DN 116:214489
 TI Preparation of (heterocyclmethyl)nitroguanidines as pesticides
 IN Minamida, Isao; Kando, Yasuyuki; Ishizuka, Hitoshi; Okauchi, Tetsuo;
 Uneme, Hideki
 PA Takeda Chemical Industries, Ltd., Japan
 SO Eur. Pat. Appl., 68 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 IC ICM C07D277-32
 ICS A01N047-44; C07D213-61
 CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 471372	A1	19920219	EP 1991-113705	19910815
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	IL 99161	A1	19961114	IL 1991-99161	19910812
	US 5256679	A	19931026	US 1991-745245	19910814
	CA 2049251	AA	19920218	CA 1991-2049251	19910815
	JP 05271207	A2	19931019	JP 1991-288208	19910815
	BR 9103540	A	19920512	BR 1991-3540	19910816
	HU 61735	A2	19930301	HU 1991-2731	19910816
	CN 1059719	A	19920325	CN 1991-105707	19910817
	CN 1028592	B	19950531		
	CN 1107836	A	19950906	CN 1994-107830	19940629

US 5696256	A 19971209	US 1995-439790	19950512
PRAI JP 1990-217356	19900817		
JP 1990-219628	19900820		
JP 1990-276628	19901015		
JP 1991-36108	19910301		
JP 1991-111987	19910516		
JP 1990-98627	19900413		
US 1991-682247	19910409		
OS MARPAT 116:214489			
GI			



AB [R1CH₂(R2)N] (R3R4N)C:NNO₂ [I; R1 = (substituted) heterocyclyl; R2 = group attached through a P or S atom, cyano, CO₂R₆, CONR₇R₈; R3 = H, (substituted) hydrocarbyl, group attached through a P or S atom, cyano, COR₉, CO₂R₁₀, CONR₁₁R₁₂; R4 = H, alkyl; R₆, R₁₀ = (substituted) hydrocarbyl, heterocyclyl; R₇, R₈, R₉, R₁₁, R₁₂ = R₆; R₇R₈N, R₁₁R₁₂N = cyclic amino group] were prep'd. Thus, 1-(2-chloro-5-thiazolylmethyl)-3,3-dimethyl-2-nitroguanidine was stirred 5 min with NaH in DMF; BrCN was added and the mixt. was stirred 1 h to give title compd. II. All I as 100 ppm sprays gave 100% kills of Nilaparvata lugens, Spodoptera litura, and Aphid gossypii.

ST nitroguanidine heterocyclylmethyl prep'n pesticide; insecticide heterocyclylmethylnitroguanidine; **thiazolylmethylnitroguanidine** prep'n insecticide

IT Insecticides

Pesticides
(heterocyclylmethyl)nitroguanidines)

IT 141205-34-7P	141205-35-8P	141205-36-9P	141205-37-0P	141205-38-1P
141205-39-2P	141205-40-5P	141205-41-6P	141205-42-7P	141205-43-8P
141205-44-9P	141205-45-0P	141205-46-1P	141205-47-2P	141205-48-3P
141205-49-4P	141205-50-7P	141205-51-8P	141205-52-9P	141205-53-0P
141205-54-1P	141205-55-2P	141205-56-3P	141205-57-4P	141205-58-5P
141205-59-6P	141205-60-9P	141205-61-0P	141205-62-1P	141205-63-2P
141205-64-3P	141205-65-4P	141205-66-5P	141205-67-6P	141205-68-7P
141205-69-8P	141205-70-1P	141205-71-2P	141205-72-3P	141205-73-4P
141205-74-5P	141205-75-6P	141205-76-7P	141205-77-8P	141205-78-9P
141205-79-0P	141205-80-3P	141205-81-4P	141205-82-5P	141205-83-6P
141205-84-7P	141227-31-8P	141227-32-9P	141227-33-0P	141227-34-1P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of, as pesticide)

IT 141205-33-6P	141205-85-8P	141205-86-9P	141205-87-0P	141205-88-1P
141205-89-2P	141205-90-5P	141205-91-6P	141227-35-2P	141227-36-3P
141227-37-4P	141227-38-5P	141227-39-6P	141227-40-9P	141227-41-0P
141227-42-1P	141227-43-2P	141227-44-3P	141227-45-4P	141227-46-5P
141227-47-6P	141227-48-7P	141227-49-8P	141227-50-1P	141227-51-2P
141227-52-3P	141227-53-4P	141227-54-5P	141227-55-6P	141227-56-7P
141227-57-8P	141227-58-9P	141227-59-0P	141227-60-3P	141227-61-4P
141227-62-5P	141227-63-6P	141227-64-7P	141227-65-8P	141227-66-9P
141227-67-0P	141227-68-1P	141227-69-2P	141227-70-5P	141227-71-6P
141227-72-7P				

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, as pesticide intermediate)

IT 74-89-5, Methylamine, reactions 79-22-1, Methyl chlorocarbonate
97-72-3, Isobutyric anhydride 98-88-4, Benzoyl chloride 103-80-0,

Phenylacetyl chloride 108-12-3 108-23-6 109-61-5 110-91-8,
 Morpholine, reactions 123-62-6 124-40-3, Dimethylamine, reactions
 124-63-0, Methanesulfonyl chloride 141-75-3, Butanoyl chloride
 501-53-1, Benzyl chloroformate 506-68-3, Cyanogen bromide 541-41-3
 543-27-1 592-34-7 594-42-3, Trichloromethanesulfonyl chloride
 638-29-9, Pentanoyl chloride 1005-56-7, Phenyl chlorothionoformate
 1710-98-1 1885-14-9, Phenyl chloroformate 2986-25-6,
 S-Methyl-N-nitroisothiourea 5271-67-0, 2-Thiophenecarbonyl chloride
 5856-79-1 10147-36-1, Propanesulfonyl chloride 24424-99-5

38870-89-2

50893-53-3, 1-Chloroethyl chloroformate 59660-22-9 120740-08-1
 131748-65-7 138149-97-0

RL: RCT (Reactant)
 (reaction of, in prepn. of pesticide)

L7 ANSWER 22 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1992:106279 CAPLUS

DN 116:106279

TI Preparation of 1-amino-1-(heterocyclylmethylamino)-2-nitro-3-hydroxy-4-halo(alkyl)-1-butenes as pesticides

IN Maienfisch, Peter; Gsell, Laurenz; Kristiansen, Odd

PA Ciba-Geigy A.-G., Switz.

SO Eur. Pat. Appl., 79 pp.

CODEN: EPXXDW

DT Patent

LA German

IC ICM C07D213-61

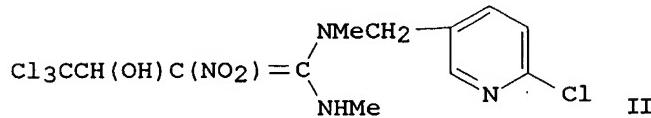
ICS A01N043-40; C07D213-38; C07D277-32; C07D213-89; A01N043-78

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))

Section cross-reference(s): 5, 27

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 453398	A2	19911023	EP 1991-810225	19910327
	EP 453398	A3	19920520		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL				
	CA 2039779	AA	19911007	CA 1991-2039779	19910404
	AU 9174091	A1	19911024	AU 1991-74091	19910404
	AU 640999	B2	19930909		
	CN 1055356	A	19911016	CN 1991-102178	19910405
	HU 56810	A2	19911028	HU 1991-1125	19910405
	BR 9101372	A	19911126	BR 1991-1372	19910405
	ZA 9102532	A	19911224	ZA 1991-2532	19910405
	JP 04234848	A2	19920824	JP 1991-100470	19910405
PRAI	CH 1990-1169		19900406		
	CH 1990-3481		19901102		
OS	MARPAT	116:106279			
GI					



AB XYZCCH(OH)(C(NO₂):C(NR₃R₄)NR₁CHR₂A [I; R₁, R₃ = H, (cyclo)alkyl; R₂ = H, alkyl; R₄ = H, (cyclo)alkyl, CH₂B; R₃R₄ = (CH₂)₄, (CH₂)₅; X = halo, haloalkyl; Y, Z = halo; A = (substituted) (arom.) (bi)heterocyclyl; B = (substituted) Ph, pyridyl, thiazolyl], were prep'd. Thus, 1-methylamino-1-[N-(6-chloropyrid-3-ylmethyl)-N-methyl]amino-2-nitroethylene was stirred with Cl₃CCHO in CH₂Cl₂ to give 94% title compds.

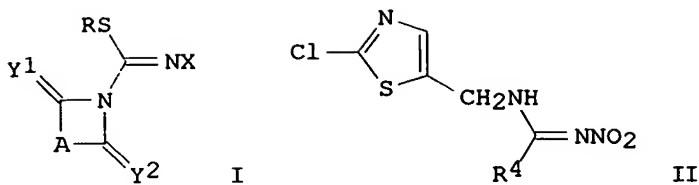
II. Numerous I at 400 ppm on rice plants gave > 80% control of

ST Nephotettix cincticeps.
 heterocyclynitroenamine prepn pesticide; insecticide
 heterocyclynitroenamine; acaricide heterocyclynitroenamine
 IT Acaricides
 Insecticides
 (heterocyclenitroenamines)
 IT 75-87-6, Chloral 375-02-0 811-96-1
 RL: RCT (Reactant)
 (condensation of, with (methylamino)nitroethylene deriv.)
 IT 120738-59-2
 RL: RCT (Reactant)
 (condensation of, with chloral, in prepn. of insecticide and
 acaricide)
 IT 133077-67-5P 137996-47-5P 137996-48-6P 137996-49-7P 137996-50-0P
 137996-51-1P 137996-52-2P 137996-53-3P 137996-54-4P 137996-55-5P
 137996-56-6P 137996-57-7P 137996-58-8P 137996-59-9P 137996-60-2P
 137996-61-3P 137996-62-4P 137996-63-5P 137996-64-6P 137996-65-7P
 137996-66-8P 137996-67-9P 137996-68-0P 137996-69-1P 137996-70-4P
 137996-71-5P 137996-72-6P 137996-73-7P 137996-74-8P 137996-75-9P
 137996-76-0P 137996-77-1P 137996-78-2P 137996-79-3P 137996-80-6P
 137996-81-7P 137996-82-8P 137996-83-9P 137996-84-0P 138023-64-0P
 138626-97-8P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except
 adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
 (Preparation); USES (Uses)
 (prepn. of, as insecticide and acaricide)

L7 ANSWER 23 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1992:21040 CAPLUS
 DN 116:21040
 TI Preparation of N-thiazolylmethyl-N"-nitroguanidines and analogs
 as acaricides and insecticides
 IN Yasuyuki, Kando; Hideki, Uneme; Isao, Minamida
 PA Takeda Chemical Industries, Ltd., Japan
 SO Eur. Pat. Appl., 37 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 IC ICM C07D207-404
 ICS C07D209-48; C07D211-88; C07D223-10; C07D213-40; C07D213-61;
 C07D277-28; C07D277-32; C07C335-40
 CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 452782	A1	19911023	EP 1991-105637	19910410
	EP 452782	B1	19970115		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	JP 05009173	A2	19930119	JP 1991-144295	19910402
	AT 147730	E	19970215	AT 1991-105637	19910410
	ES 2096598	T3	19970316	ES 1991-105637	19910410
	CA 2040385	AA	19911014	CA 1991-2040385	19910412
	HU 58049	A2	19920128	HU 1991-1227	19910412
	IL 97839	A1	19970218	IL 1991-97839	19910412
	US 5696256	A	19971209	US 1995-439790	19950512
PRAI	JP 1990-98627	19900413			
	JP 1990-217356	19900817			
	US 1991-682247	19910409			
OS	MARPAT 116:21040				
GI					



AB R₁R₂NC(:NNO₂)NR₃(CH₂)_nB [B = (un)substituted (hetero)cyclic group; R₁, R₃ = H, acyl, (un)substituted hydrocarbyl; R₂ = H, (un)substituted hydrocarbyl; NR₁R₂ = cyclic amino group; n = 0, 1] were prep'd. as insecticides and acaricides (no data) together with their intermediates I [R = acyl, (un)substituted hydrocarbyl; A = (un)substituted hydrocarbyldiyl; X = electron-withdrawing group; Y₁, Y₂ = O, S]. Thus, MeSC(:NNO₂)NH₂ was cyclocondensed with phthaloyl chloride to give I (A = 1,2-phenylenediyl, R = Me, X = NO₂, Y₁ = Y₂ = O) which was condensed with 2-chloro-5-(aminomethyl)thiazole to give isothiourea II (R₄ = SME). The latter was condensed with MeNH₂ to give title compd. II (R₄ = MeNH).

ST nitroguanidine **thiazolylmethyl** prepn acaricide insecticide

IT Acaricides

Insecticides

(N-thiazolylmethyl-N''-nitroguanidines and analogs)

IT 59660-22-9P 132219-48-8P 137589-64-1P 138149-96-9P 138149-97-0P
138149-98-1P 138149-99-2P 138150-00-2P 138150-01-3P 138150-02-4P
138150-03-5P 138150-04-6P 138150-05-7P 138169-89-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. and reaction of, in prepn. of insecticides and acaricides)

IT 35089-65-7P 131748-47-5P 131748-48-6P 131748-49-7P 131748-50-0P
131748-51-1P 131748-53-3P 131748-54-4P 131748-55-5P 131748-56-6P
131748-57-7P 131748-58-8P 131748-59-9P 131748-60-2P 131748-61-3P
131748-65-7P 131748-66-8P 131748-69-1P 131748-70-4P 131748-71-5P
131748-72-6P 131748-73-7P 131748-74-8P 131748-75-9P 131748-76-0P
131748-85-1P 131748-86-2P 131768-12-2P 131768-13-3P 135018-15-4P
137589-63-0P 138149-87-8P 138149-88-9P 138149-89-0P 138149-90-3P
138149-91-4P 138149-92-5P 138149-93-6P 138149-94-7P 138149-95-8P
138150-06-8P 138150-07-9P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of, as insecticide and acaricide)

IT 88-95-9, Phthaloyl chloride 100-46-9, Benzylamine, reactions

111-50-2,

Adipic acid dichloride 543-20-4, Succinic acid chloride 2873-74-7,
Pantanediol dichloride 2986-25-6, S-Methyl-N-nitroisothiourea ...
21062-20-4, Diglycolic acid dichloride 60901-05-5 97004-04-1,
(6-Chloro-3-pyridyl)methylamine 120739-77-7 120740-08-1,
2-Chloro-5-(aminomethyl)thiazole

RL: RCT (Reactant)

(reaction of, in prepn. of insecticides and acaricides)

L7 ANSWER 24 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1991:680003 CAPLUS

DN 115:280003

TI Preparation of (heteroaryl methyl)nitroguanidines as insecticides

IN Nanjo, Katsumi; Takasuka, Kiyoshi; Segami, Shigenori; Kariya, Akinori

PA Agro-Kanesho Co., Ltd., Japan

SO Eur. Pat. Appl., 15 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-61

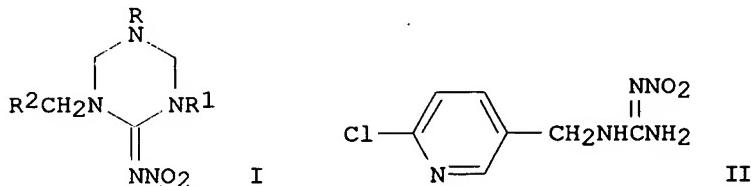
ICS C07D277-32; A01N047-44

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 425978	A2	19910508	EP 1990-120294	19901023
	EP 425978	A3	19911106		
	R: CH, DE, FR, GB, IT, LI				
	JP 03200768	A2	19910902	JP 1989-328888	19891219
	AU 9063964	A1	19910530	AU 1990-63964	19901010
	US 5166164	A	19921124	US 1990-596039	19901011
	CN 1051728	A	19910529	CN 1990-109217	19901023
PRAI	JP 1989-276633	19891024			
	JP 1989-328888	19891219			
OS	MARPAT 115:280003				
AB	XCH2NRC(:NNO2)NR1R2 (I; R1-R2 = H, Me; X = 6-chloro-3-pyridinyl, 2-chloro-5-thiazolyl) were prep'd. by reaction of MeSC(:NNO2)NR1R2 with XCH2NHR (R = H, Me; X as above), optionally followed				
	by methylation of the resulting nitroguanidine. Thus, a mixt. of 1.50 g 2-chloro-N-methyl-5-thiazolemethanamine and 1.13 g MeSC(:NNO2)NH2 in 6 mL EtOH was refluxed 6 h to give 1.00 g title compd. (I; R = Me, R1 = R2 = H, X = 2-chloro-5-thiazolyl) which at 500 ppm on rice seedlings gave 100% kill of second instar larvae of green rice leafhopper which had acquired resistance to chems., vs. 70% kill at the same concn. for (MeO)2P(:S)OC6H3(Me)NO2-3,4 (sumithion) as a control.				
ST	chlorothiazolylnitroguanidine prepn insecticide; thiazolylmethanamine substitution methylnitroisothiourea				
IT	Insecticides (heteroarylmethylnitroguanidines)				
IT	120739-62-0 120740-06-9 120740-08-1				
RL: RCT (Reactant)	(condensation reaction of, with nitroisothiourea deriv., in prepn. of insecticide)				
IT	2986-25-6, S-Methyl-N-nitroisothiourea 59660-22-9				
RL: RCT (Reactant)	(condensation reaction of, with pyridine- and thiazolemethanamine derivs., in prepn. of insecticides)				
IT	131748-47-5P 131748-49-7P 131748-55-5P 131748-56-6P 131748-59-9P 131748-70-4P 131748-77-1P 131768-12-2P 135018-15-4P				
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)	(prepn. of, as insecticide)				
L7	ANSWER 25 OF 33 CAPLUS COPYRIGHT 1999 ACS				
AN	1991:583374 CAPLUS				
DN	115:183374				
TI	Preparation of hexahydrotriazine compounds as insecticides				
IN	Wu, Frank; Kariya, Akinori; Katsuyama, Noriyoshi; Tsuji, Atsushi; Takasuka, Kiyoshi; Segami, Shigenori; Nanjo, Katsumi; Sato, Junko				
PA	Agro-Kanesho Co., Ltd., Japan				
SO	Eur. Pat. Appl., 16 pp.				
	CODEN: EPXXDW				
DT	Patent				
LA	English				
IC	ICM C07D251-04				
	ICS A01N043-64				
CC	28-19 (Heterocyclic Compounds (More Than One Hetero Atom))				
	Section cross-reference(s): 5				
FAN.CNT 1					
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 428941	A1	19910529	EP 1990-121383	19901108

EP 428941	B1	19950503		
R: CH, DE, FR, GB, IT, LI				
JP 03218370	A2	19910925	JP 1990-24199	19900202
JP 06000776	B4	19940105		
AU 9065623	A1	19910516	AU 1990-65623	19901031
AU 628229	B2	19920910		
CN 1060656	A	19920429	CN 1990-109039	19901110
CN 1026648	B	19941123		
CN 1098719	A	19950215	CN 1994-103331	19940325
PRAI	JP 1989-292675	19891110		
	JP 1990-24199	19900202		
OS	MARPAT	115:183374		
GI				



AB Hexahydrotriazine derivs. [I; R = alkyl, alkenyl; R1 = H, alkyl, alkenyl, alkynyl, (6-chloro-3-pyridyl)methyl; R2 = 6-chloro-3-pyridyl, 2-chlorothiazol-5-yl] are prep'd. A soln. of Et₃N in THF was added to a suspension of 0.6 g guanidine deriv. II and 0.4 g (ClCH₂)₂NMe in THF with stirring under cooling to give 0.78 g I (R = Me, R1 = H, R2 = 6-chloro-3-pyridyl), which killed 100% green rice leafhopper larvae at 500 ppm. Also prep'd. and tested were 11 addnl. I.

ST triazine prepn insecticide

IT Insecticides

(hexahydrotriazine compds.)

IT 131748-56-6

RL: RCT (Reactant)

(cyclocondensation of, with amine derivs., in prepn. of insecticide)

IT 75-04-7, Ethylamine, reactions 34645-08-4

RL: RCT (Reactant)

(cyclocondensation of, with nitroguanidine deriv., in prepn. of insecticide)

IT 133258-61-4P 133258-66-9P 136516-17-1P 136516-18-2P 136516-19-3P
136516-20-6P 136516-21-7P 136516-22-8P 136516-23-9P 136516-24-0P
136516-25-1P 136544-01-9P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of, as insecticide)

IT 136516-16-0

RL: RCT (Reactant)

(reaction of, with chloropyridylmethyl chloride, in prepn. of insecticide)

IT 70258-18-3 105827-91-6, 2-Chloro-5-thiazolylmethyl chloride

RL: RCT (Reactant)

(reaction of, with triazine deriv., in prepn. of insecticide)

L7 ANSWER 26 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1991:471585 CAPLUS

DN 115:71585

TI Guanidine derivatives as insecticides

IN Kristiansen, Odd; Maienfisch, Peter; Gsell, Laurenz

PA Ciba-Geigy A.-G., Switz.

SO Eur. Pat. Appl., 48 pp.

CODEN: EPXXDW

DT Patent

LA German

IC ICM C07D213-61

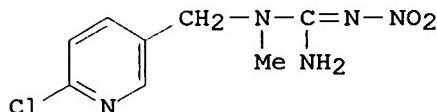
ICS C07D213-40; A01N043-40; C07D277-32; C07D401-12

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))

Section cross-reference(s): 5, 27

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 418199	A2	19910320	EP 1990-810668	19900904
	EP 418199	A3	19910612		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL				
	CA 2025072	AA	19910314	CA 1990-2025072	19900911
	DD 297639	A5	19920116	DD 1990-343952	19900911
	AU 9062456	A1	19910321	AU 1990-62456	19900912
	CN 1050186	A	19910327	CN 1990-107654	19900912
	HU 54872	A2	19910429	HU 1990-5882	19900912
	ZA 9007265	A	19910626	ZA 1990-7265	19900912
	BR 9004550	A	19910910	BR 1990-4550	19900912
	JP 03109374	A2	19910509	JP 1990-243700	19900913
PRAI	CH 1989-3335		19890913		
	CH 1990-1078		19900402		
OS	MARPAT 115:71585				
GI					



AB ACHR2NR1C(:NNO2)NR3R4 [R1, R3 = H, C1-4 alkyl, C3-6 cycloalkyl; R2 = H, C1-4 alkyl; R4 = H, C1-4 alkyl, C3-6 cycloalkyl, CHR5B; R3R4 = (CH2)4, (CH2)5; R5 = H, C1-4 alkyl; A = (un)substituted, arom. or (un)satd., (bi)cyclic (hetero)ring; B = (un)substituted Ph, (un)substituted 3-pyridyl, (un)substituted 5-thiazolyl] and their salts (with proviso), were prep'd., e.g., by reaction of amines HNR1CHR2A with nitroisothioureas MeSC(:NNO2)NR3R4. Thus, a mixt. of N-methyl-(2-chloropyrid-5-yl)methylamine, S-methyl-N-nitroisothiourea, and KHSO4 in EtOH was refluxed 4.5 h to give title compd. (I) which at 400 ppm on rice plants gave >80% redn. of the population of Nilaparvata lugens.

ST nitroguanidine prepn insecticide

IT Insecticides

(nitroguanidines)

IT 104-83-6, 4-Chlorobenzyl chloride

RL: RCT (Reactant)

(benzylation by, of nitroguanidine deriv., in prepn. of insecticide)

IT	35089-65-7P	131748-47-5P	131748-48-6P	131748-49-7P	131748-50-0P
	131748-53-3P	131748-54-4P	131748-56-6P	131748-57-7P	131748-59-9P
	131748-65-7P	131748-70-4P	131768-12-2P	133001-33-9P	135018-02-9P
	135018-03-0P	135018-04-1P	135018-05-2P	135018-06-3P	135018-07-4P
	135018-08-5P	135018-09-6P	135018-10-9P	135018-11-0P	135018-12-1P
	135018-13-2P	135018-14-3P	135018-15-4P	135018-16-5P	135018-17-6P
	135018-18-7P	135018-19-8P	135018-20-1P	135051-39-7P	

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of, as insecticide)

IT 41288-91-9

RL: RCT (Reactant)

(reaction of, with isothiourea deriv., in prepn. of insecticide)

IT 120739-84-6

RL: RCT (Reactant)

(reaction of, with nitroguanidine, in prepn. of insecticide)

IT 556-88-7, Nitroguanidine 2986-25-6, S-Methyl-N-nitroisothiourea

RL: RCT (Reactant)

(reaction of, with pyridylmethylamine deriv., in prepn. of insecticide)

L7 ANSWER 27 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1991:185533 CAPLUS

DN 114:185533

TI Preparation of nitro-substituted heterocyclic compounds as insecticides

IN Shiokawa, Kozo; Tsuboi, Shinichi; Moriya, Koichi; Hattori, Yumi; Honda, Ikuro; Shibuya, Katsuhiko

PA Nihon Tokushu Noyaku Seizo K. K., Japan

SO Eur. Pat. Appl., 54 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-74

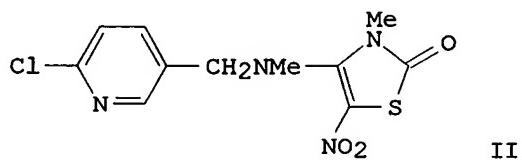
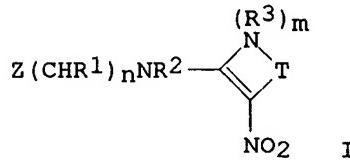
ICS A01N043-30; A01N043-54; A01N043-78; C07D401-12; C07D239-12;
C07D417-12

CC 28-16 (Heterocyclic Compounds (More Than One Hetero Atom))

Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 398084	A2	19901122	EP 1990-108414	19900504
	EP 398084	A3	19920429		
	R: BE, CH, DE, FR, GB, IT, LI, NL				
	JP 03197460	A2	19910828	JP 1990-25190	19900206
	US 5081132	A	19920114	US 1990-518684	19900503
	EP 606105	A1	19940713	EP 1994-101886	19900504
	EP 606105	B1	19981125		
	R: BE, CH, DE, FR, GB, IT, LI, NL				
	US 5219869	A	19930615	US 1992-943950	19920911
	US 5314897	A	19940524	US 1993-90566	19930712
PRAI	JP 1989-121366	19890517			
	US 1990-518684	19900503			
	EP 1990-108414	19900504			
	US 1991-760412	19910916			
	US 1992-898064	19920612			
OS	MARPAT 114:185533				
GI					



AB Title compds. I [R¹ = H, cyano, C1-4 alkyl; R² = H, (substituted) C1-4 alkyl, halo, C1-4 alkyl, halo, C1-4 alkoxy, C3-4 alkenyl, C3-4 alkynyl, (substituted) Ph or PhCH₂, methylated (substituted) heterocyclyl; T = (substituted) 2- or 3-membered divalent or trivalent chain comprising hetero and(or) C atoms; Z = (substituted) Ph or heterocyclyl; m, n = 0] are prep'd. A mixt. of 3-methyl-4-(methylthio)-5-nitro-2-thiazolone (prepn. given), N-(2-chloro-5-(pyridylmethyl)-N-methylamine and EtOH was refluxed for 5 h and cooled to room temp. to give

the title heterocycle II. II at 200 ppm gave 100% mortality of Nephrotettix cincticeps on rice plant.

ST pyridylmethylaminoheterocycle prepn insecticide; **thiazolone**
pyridylmethylamino insecticide; heterocycle nitro substituted
insecticide;
pyrimidine pyridylmethylamino insecticide

IT Insecticides
(nitrosubstituted heterocyclic compds.)

IT 129950-36-3P 133287-11-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. and reaction of, in prepn. of nitro-substituted heterocyclic
insecticides)

IT 129438-21-7P 129438-30-8P 129438-31-9P 129438-33-1P 129438-35-3P
129438-39-7P 129438-58-0P 129452-08-0P 129950-09-0P 129950-10-3P
129950-15-8P 133286-90-5P 133286-91-6P 133286-92-7P 133286-93-8P
133286-94-9P 133286-95-0P 133286-96-1P 133286-97-2P 133286-98-3P
133286-99-4P 133287-00-0P 133287-01-1P 133287-02-2P 133287-03-3P
133287-04-4P 133287-05-5P 133287-06-6P 133287-07-7P 133287-08-8P
133287-09-9P 133287-10-2P 133304-00-4P
RL: AGR (Agricultural use); BAC (Biological activity or effector, except
adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
(Preparation); USES (Uses)
(prepn. of, as insecticide)

IT 50-00-0, Formaldehyde, reactions 74-89-5, Methylamine, reactions
2757-23-5 5470-18-8, 2-Chloro-3-nitropyridine 6542-88-7, Aminoacetal
13623-94-4, 2,2-Bis(methylthio)-1-nitroethylene 61832-41-5

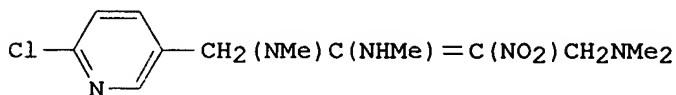
120738-58-1
120739-62-0
RL: RCT (Reactant)
(reaction of, in prepn. of nitro-substituted heterocyclic
insecticides)

L7 ANSWER 28 OF 33 CAPLUS COPYRIGHT 1999 ACS
AN 1991:164015 CAPLUS
DN **114:164015**
TI Preparations of (pyridylalkyl)diaminoethylenes as insecticides
IN Uneme, Hideki; Minamida, Isao; Okauchi, Tetsuo
PA Takeda Chemical Industries, Ltd., Japan
SO Eur. Pat. Appl., 23 pp.
CODEN: EPXXDW
DT Patent
LA English
IC ICM C07D213-61
ICS C07D277-32; A01N043-40; A01N043-78; C07D213-643; C07D213-74;
C07D213-70; C07D213-38
CC 27-16 (Heterocyclic Compounds (One Hetero Atom))
Section cross-reference(s): 15

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 392560	A2	19901017	EP 1990-107120	19900412
	EP 392560	A3	19920108		
	EP 392560	B1	19951227		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	IL 94027	A1	19950315	IL 1990-94027	19900406
	BR 9001734	A	19910604	BR 1990-1734	19900411
	JP 03169861	A2	19910723	JP 1990-97363	19900411
	IN 170550	A	19920411	IN 1990-MA269	19900411
	CA 2014490	AA	19901014	CA 1990-2014490	19900412
	US 5438065	A	19950801	US 1990-507776	19900412
	AT 132139	E	19960115	AT 1990-107120	19900412
	ES 2081314	T3	19960301	ES 1990-107120	19900412
	HU 53780	A2	19901228	HU 1990-2438	19900413
	HU 207202	B	19930329		
	CN 1046896	A	19901114	CN 1990-102111	19900414

CN 1036112 B 19971015
PRAI JP 1989-95580 19890414
JP 1989-201980 19890802
OS MARPAT 114:164015
GI



I

AB Title compds. R1(CH2)nNR2C(NR3R4):CXCHR5Y (R1 = (substituted) heterocycl; R2, R3, R4 = H, (substituted) hydrocarbyl, R3R4N = heterocycl; R5 = H, (substituted) hydrocarbyl, (substituted) heterocycl; X = electron attractant; Y = R6O, R6 = H, (substituted) hydrocarbyl, -heterocycl, (substituted) amino), etc.; n = 0, 1) or a salt thereof, are prep'd.
1-[N-(6-Chloro-3-pyridylmethyl)-N-methylamino]-1-(methylamino)-2-nitroethylene, aq. CH2O, aq. Me2NH and MeCN were stirred at room temp. for 8.5 h to give the pyridine deriv. I. I at 500 and 100 ppm resulted in 100% mortality against Nilaparvata lugens and Aphis gossypii, resp.
ST pyridylmethyldiaminoethylene substituted prep'n insecticide; thiazolylmethyldiaminoethylene substituted prep'n insecticide; insecticide heterocyclidiaminoethylene substituted
IT Insecticides (substituted heterocyclidiaminoethylene)
IT 133077-59-5P 133077-60-8P 133077-61-9P 133077-62-0P 133077-63-1P
133077-64-2P 133077-65-3P 133077-66-4P 133077-67-5P 133077-68-6P
133077-69-7P 133077-70-0P 133077-71-1P 133077-72-2P 133077-73-3P
133077-74-4P 133077-75-5P 133077-76-6P 133077-77-7P 133077-78-8P
133077-79-9P 133077-80-2P 133077-81-3P 133077-82-4P 133077-83-5P
133077-84-6P 133077-85-7P 133077-86-8P 133077-87-9P 133077-88-0P
133077-89-1P 133077-90-4P 133077-91-5P 133105-15-4P
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of, as insecticide)
IT 50-00-0, Formaldehyde, reactions 106-54-7, p-Chlorothiophenol
302-17-0, Chloral hydrate 120738-59-2
RL: RCT (Reactant)
(reaction of, in prepn. of insecticides)

L7 ANSWER 29 OF 33 CAPLUS COPYRIGHT 1999 ACS.

AN 1991:61936 CAPLUS

DN 114:61936

TI Preparation of pyridine derivatives and other heterocycles as insecticides

IN Aoki, Isao; Tabuchi, Takanori; Minamida, Isao

PA Takeda Chemical Industries, Ltd., Japan

SO Eur. Pat. Appl., 34 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-36

ICS C07D213-61; C07D213-64; C07D215-12; C07D241-12; C07D277-28;
C07D277-32; C07C205-08

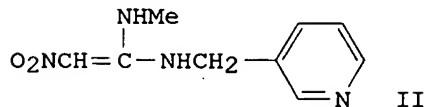
CC 27-16 (Heterocyclic Compounds (One Hetero Atom))

Section cross-reference(s): 5

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	EP 381130	A2	19900808	EP 1990-101778	19900130
	EP 381130	A3	19901031		
	R: CH, DE, FR, GB, IT, LI				
	IL 93159	A1	19961114	IL 1990-93159	19900124
	HU 52751	A2	19900828	HU 1990-609	19900131
	HU 209439	B	19940628		
	JP 02275841	A2	19901109	JP 1990-22503	19900131
	US 5364989	A	19941115	US 1992-908725	19920706
PRAI	JP 1989-23356	19890131			
	US 1990-473173	19900131			
	US 1990-527898	19900524			
OS	MARPAT 114:61936				
GI					



AB The title compds. O₂NCH:C(NR₂R₃)NR₁CnH₂nA [R₁ = H, alkyl, haloalkyl, aralkyl, etc.; R₂ = H, alkyl, aralkyl; R₃ = H, alkyl, haloalkyl, hydroxyalkyl, etc.; A = (substituted) 3- or 4-pyridyl, pyrazinyl, 2-, 4-, or 5-thiazolyl, etc.; n = 0-2] were prep'd. by reaction of X₁X₂X₃CCH₂NO₂ (I; X₁, X₂ = F, Cl, Br, iodo; X₃ = Cl, Br) with R₁NHCnH₂nA. I were prep'd. by reactive of X₁X₂C:CH₂ with HNO₃ or its salts and HCl or HBr or its salt. N-(6-Chloro-3-pyridyl)methyl-N-ethylamine was added to

a mixt. of C₁₃CCH₂NO₂ and K₂CO₃ in MeCN. After addn. of K₂CO₃ and MeNH₂ in MeOH, the reaction mixt. was stirred for 1 h at 18-20.degree. to give 62.7% 1-[N-6-chloro-3-pyridyl)methyl-N-ethyl]amino-1-methylamino-2-nitroethylene. Pyridine deriv. II at 500 ppm gave 100% kill of Nilaparvata lugens larvae.

ST pyridine substituted prepn insecticide; heterocyclic compd prepn insecticide; insecticide substituted pyridine heterocyclic compd

IT Insecticides

(pyridine derivs. and other heterocycles)

IT 630-20-6P, 1,1,1,2-Tetrachloroethane

RL: FORM (Formation, nonpreparative); SPN (Synthetic preparation); PREP (Preparation)

(formation of, in prepn. of insecticide)

IT 7697-37-2, Nitric acid, reactions

RL: RCT (Reactant)

(nitration by, of dichloroethylene)

IT 75-35-4, 1,1-Dichloroethylene, reactions

RL: RCT (Reactant)

(nitration of)

IT 6061-04-7P, 1,1-Dichloro-2-nitroethylene

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)

(prepn. and reaction of, in prepn. of insecticide)

IT	90787-25-0P	120738-35-4P	120738-36-5P	120738-37-6P	120738-38-7P
	120738-39-8P	120738-40-1P	120738-41-2P	120738-42-3P	120738-43-4P
	120738-44-5P	120738-45-6P	120738-46-7P	120738-47-8P	120738-48-9P
	120738-49-0P	120738-50-3P	120738-51-4P	120738-52-5P	120738-53-6P
	120738-54-7P	120738-55-8P	120738-56-9P	120738-58-1P	120738-59-2P
	120738-61-6P	120738-62-7P	120738-63-8P	120738-64-9P	120738-65-0P
	120738-66-1P	120738-67-2P	120738-68-3P	120738-69-4P	120738-70-7P
	120738-71-8P	120738-72-9P	120738-73-0P	120738-74-1P	120738-75-2P
	120738-76-3P	120738-77-4P	120738-78-5P	120738-79-6P	120738-80-9P
	120738-81-0P	120738-82-1P	120738-85-4P	120738-86-5P	120738-87-6P
	120738-88-7P	120738-89-8P	120738-90-1P	120738-91-2P	120738-92-3P
	120738-93-4P	120738-96-7P	120738-97-8P	120738-98-9P	120738-99-0P
	120739-00-6P	120739-03-9P	120739-04-0P	120739-05-1P	120739-06-2P

120739-07-3P	120739-08-4P	120739-09-5P	120739-10-8P	120739-11-9P
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120739-18-6P	120739-19-7P	120739-20-0P	120739-21-1P	120739-22-2P
120739-23-3P	120739-24-4P	120739-25-5P	120739-26-6P	120739-27-7P
120739-28-8P	120739-29-9P	120739-30-2P	120739-31-3P	120739-32-4P
120739-33-5P	120739-34-6P	120739-35-7P	120739-36-8P	120739-46-0P
120739-55-1P	120739-57-3P	120739-59-5P	120770-85-6P	120770-86-7P
120770-87-8P	131427-96-8P	131427-97-9P	131427-98-0P	131427-99-1P
131428-00-7P	131428-01-8P	131428-02-9P	131428-03-0P	131428-04-1P
131428-05-2P	131428-06-3P	131428-07-4P	131428-08-5P	

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of, as insecticide)

IT 64326-81-4P, 1,1,1-Trichloro-2-nitroethane 97987-61-6P

RL: SPN (Synthetic preparation); PREP (Preparation)

(prepn. of, as insecticide intermediate)

IT 97004-04-1 120739-77-7 120740-08-1

RL: RCT (Reactant)

(reaction of, in prepn. of insecticide)

L7 ANSWER 30 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1990:532013 CAPLUS

DN 113:132013

TI Preparation of (pyridylmethyl)-containing cyano compounds and analogs as insecticides

IN Shiokawa, Kozo; Tsuboi, Shinichi; Moriya, Koichi; Honda, Ikuro; Hattori, Yumi; Shibuya, Katsuhiko

PA Nihon Tokushu Noyaku Seizo K. K., Japan

SO Eur. Pat. Appl., 56 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM C07D213-61

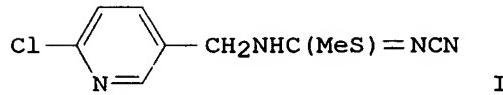
ICS A01N047-44; C07D213-75; C07D239-26; C07D277-32; A01N047-42

CC 27-16 (Heterocyclic Compounds (One Hetero Atom))

Section cross-reference(s): 5, 28

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 364844	A1	19900425	EP 1989-118689	19891007
	EP 364844	B1	19940316		
	R: BE, CH, DE, FR, GB, IT, LI, NL				
	JP 02209868	A2	19900821	JP 1989-57813	19890313
	JP 2884412	B2	19990419		
	US 5066808	A	19911119	US 1990-584398	19900914
	US 5384324	A	19950124	US 1993-55402	19930430
PRAI	JP 1988-264020	19881021			
	JP 1989-57813	19890313			
	US 1989-419428	19891010			
	US 1990-584398	19900914			
	US 1991-678382	19910401			
OS	MARPAT	113:132013			
GI					



AB Title compds. Z(CHR1)mNR2CR3:NCN [Z = halopyridyl and other 5- and 6-membered heterocyclyl; R1 = H, cyano, C1-4 alkyl; R2 = H, C1-6 alkyl,

(un)substituted C3-4 alkenyl, C3-4 alkynyl, (un)substituted C3-8 cycloalkyl, (substituted) Ph, (substituted) PhCH₂, HO, Cl-4 alkoxy, ZCH₂; R₃ = R₄₀, R_{4S}, R₅R₆N; R₄ = Cl-6 alkyl, C3-4 alkenyl, C3-4 alkynyl, C3-8 cycloalkyl, etc.; R₅, R₆ = H, (un)substituted C1-9 alkyl, (alkyl)amino, halo, OH, SH, alkoxy, etc., or R₅R₆N = 3-7-membered heterocyclyl; m = 0, 1] are prep'd. 5-(Aminomethyl)-2-chloropyridine and (MeS)2C:NCN reacted

to

give the pyridine compd. I. I at 50 ppm showed 100% mortality against Nephrotettix cincticeps and Nilaparvata lugens.

ST pyridylmethyl cyano compd prep insecticide; pyrimidinylmethyl cyano compd

prep insecticide; thiazolylmethyl cyano compd prep insecticide; heterocyclylmethyl cyano compd prep insecticide

IT Insecticides

(heterocyclylmethyl cyano compds.)

IT	129478-21-3P	129478-22-4P	129478-23-5P	129478-24-6P	129478-25-7P
	129478-26-8P	129478-27-9P	129478-28-0P	129478-29-1P	129478-30-4P
	129478-31-5P	129478-32-6P	129478-33-7P	129478-34-8P	129478-35-9P
	129478-36-0P	129478-37-1P	129478-38-2P	129478-39-3P	129478-40-6P
	129478-41-7P	129478-42-8P	129478-43-9P		

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prep'n. of, as insecticide)

IT 5848-24-8 10191-60-3 15760-26-6 97004-04-1, 5-(Aminomethyl)-2-chloropyridine 105827-91-6, 2-Chloro-5-(chloromethyl)thiazole

RL: RCT (Reactant)

(reaction of, in prep'n. of insecticides)

L7 ANSWER 31 OF 33 CAPLUS COPYRIGHT 1999 ACS

AN 1990:158237 CAPLUS

DN 112:158237

TI Preparation and formulation of thiazole-5-carboxylic acid amides as herbicide antidotes

IN Nyffeler, Andreas; Toepfl, Werner

PA Ciba-Geigy A.-G., Switz.

SO Eur. Pat. Appl., 29 pp.

CODEN: EPXXDW

DT Patent

LA German

IC ICM C07D277-56

ICS A01N025-32

CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))

Section cross-reference(s): 5

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 335831	A1	19891004	EP 1989-810171	19890307
	R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	RO 103451	B1	19921115	RO 1989-138628	19890310
	FI 8901180	A	19890916	FI 1989-1180	19890313
	DD 283537	A5	19901017	DD 1989-326547	19890313
	PL 156391	B1	19920331	PL 1989-278186	19890313
	DK 8901231	A	19890916	DK 1989-1231	19890314
	NO 8901098	A	19890918	NO 1989-1098	19890314
	AU 8931291	A1	19890921	AU 1989-31291	19890314
	AU 608700	B2	19910411		
	CN 1035753	A	19890927	CN 1989-101360	19890314
	ZA 8901920	A	19891129	ZA 1989-1920	19890314
	JP 02015004	A2	19900118	JP 1989-61951	19890314
	HU 50266	A2	19900129	HU 1989-1225	19890314
	SU 1797459	A3	19930223	SU 1989-4613647	19890314
	BR 8901194	A	19891031	BR 1989-1194	19890315
PRAI	CH 1988-968		19880315		
	CH 1988-4673		19881219		

OS MARPAT 112:158237
 GI For diagram(s), see printed CA Issue.
 AB Title amides I [A = NR1R2, NR3NR4(CO)mR5; R1-R5 = H, (un)substituted (cyclo)alkyl, (cyclo)alkenyl, alkynyl, EmU, EmQ; NR1R2, NR4R5 = (un)substituted heterocyclyl; E = (un)substituted (un)interrupted alkylene
 or alkenylene; U = (un)substituted Ph or naphthyl; Q = heterocyclyl; m = 0, 1] were prep'd. as herbicide antidotes for sulfonylureas, haloacetanilides, and aryloxyphenoxypropionates. Thus, amidation of 2-chloro-4-trifluoromethylthiazole-5-carboxylic acid chloride with 1-cyano-1-methylaminocyclopentane in MeCN contg. Et3N gave 83% I (A = NR1R2, R1 = Me, R2 = 1-cyanocyclopent-1-yl) (II). At 8000 g/ha preemergence with an equal rate of metolachlor, II gave 70% protection of corn and no protection of Echinochloa crus-galli. A list of I, various formulations, and addnl. biol. data are given.

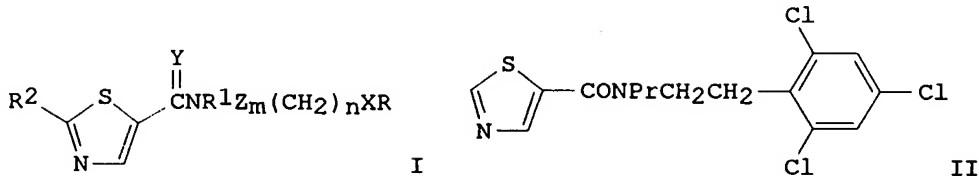
ST thiazolecarboxamide prepn herbicide antidote
 IT Herbicide antidotes
 (thiazolecarboxamide derivs.)
 IT 72850-62-5
 RL: RCT (Reactant)
 (amidation of)
 IT 55793-49-2
 RL: RCT (Reactant)
 (amidation of, with thiazolecarbonyl chloride deriv.)
 IT 5329-12-4, 2,4,6-Trichlorophenylhydrazine
 RL: RCT (Reactant)
 (condensation of, with thiazolecarbonyl chloride deriv.)
 IT 15972-60-8 51218-45-2, Metolachlor 51218-49-6, Pretilachlor
 75942-79-9, Trimexachlor 92080-03-0 126048-63-3
 RL: RCT (Reactant)
 (herbicide antidotes for)
 IT 72850-84-1P 117413-56-6P 117545-74-1P 117546-07-3P 117546-08-4P
 117546-09-5P 117546-10-8P 117546-11-9P 117546-12-0P 117546-13-1P
 117546-14-2P 117546-15-3P 117546-16-4P 117546-18-6P 117546-19-7P
 117546-20-0P 117546-21-1P 117546-22-2P 117546-23-3P 117546-26-6P
 117546-27-7P 117546-29-9P 117546-30-2P 117546-31-3P 117546-32-4P
 117546-34-6P 117546-35-7P 117546-36-8P 117546-37-9P 117546-38-0P
 117546-39-1P 117546-40-4P 117546-41-5P 117546-42-6P 117546-43-7P
 117546-44-8P 117546-45-9P 126048-56-4P 126048-57-5P 126048-58-6P
 126048-59-7P 126048-60-0P 126048-61-1P 126048-62-2P 126069-48-5P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of, as herbicide antidote)

L7 ANSWER 32 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1989:407393 CAPLUS
 DN 111:7393
 TI Preparation of 5-thiazolecarboxamides as plant fungicides
 IN Wilson, John Robert Howe; Haddock, Ernest
 PA Shell Internationale Research Maatschappij B. V., Neth.
 SO Eur. Pat. Appl., 18 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 IC ICM C07D277-56
 ICS A01N043-78
 CC 28-7 (Heterocyclic Compounds (More Than One Hetero Atom))
 Section cross-reference(s): 5
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 296673	A1	19881228	EP 1988-201227	19880615
	EP 296673	B1	19940309		

R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE

US 4877802	A	19891031	US 1988-199430	19880527
CA 1328867	A1	19940426	CA 1988-568279	19880601
AT 102611	E	19940315	AT 1988-201227	19880615
ES 2061628	T3	19941216	ES 1988-201227	19880615
DK 8803467	A	19881226	DK 1988-3467	19880623
DK 169743	B1	19950213		
AU 8818329	A1	19890105	AU 1988-18329	19880623
AU 615658	B2	19911010		
JP 01026572	A2	19890127	JP 1988-153706	19880623
CN 1033626	A	19890705	CN 1988-103766	19880623
CN 1022015	B	19930908		
SU 1579458	A3	19900715	SU 1988-4355953	19880623
PRAI GB 1987-14920		19870625		
EP 1988-201227		19880615		
OS MARPAT 111:7393				
GI				



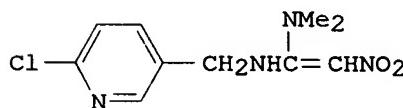
AB The title compds. [I; R = (un)substituted aryl; R1 = H, (un)substituted alkyl, alkenyl, alkynyl; R2 = H, halo, (halo)alkyl, (halo)alkoxy, alkylthio, OH, cyano, NO₂, NH₂, (di)alkylamino, morpholino; X = O, S, CO, R₄R₅C; R₄, R₅ = H, alkoxy; Y = O, S; Z = C₆H₄; n = 0, 6; m = 0, 1] and their acid salts and metal complexes were prep'd. as fungicides for plant protection. Me 2-amino-5-thiazolecarboxylate (79 g) was refluxed with Me(CH₂)₄ONO in dioxane to give 32.0 g Me 5-thiazolecarboxylate which (15 g) was saponified to give 12.1 g free acid. The latter (4.8 g) was converted to its acid chloride and treated with 10.7 g N-[2-(2,4,6-trichlorophenoxy)ethyl] propylamine (prepn. given) in pyridine to give 8.6 g title compd. II. At 1 kg/ha II gave >80% control of, e.g., Leptosphaeria nodorum on wheat and Pyricularia oryzae on rice.

ST thiazolecarboxamide prep'n agrochem fungicide
 IT Fungicides and Fungistats
 (agrochem., (phenylalkyl)thiazolecarboxamides)
 IT 107-10-8, Propylamine, reactions
 RL: RCT (Reactant)
 (alkylation of, by (bromoethoxy)trichlorobenzene)
 IT 110-78-1, Propyl isocyanate
 RL: RCT (Reactant)
 (carbamoylation of, of chlorothiazole)
 IT 3034-52-4, 2-Chlorothiazole
 RL: RCT (Reactant)
 (carbamoylation of, by Pr isocyanate)
 IT 6633-61-0, Methyl 2-amino-5-thiazolecarboxylate
 RL: RCT (Reactant)
 (diazotization and deamination of)
 IT 98-60-2, 4-Chlorobenzenesulfonyl chloride
 RL: RCT (Reactant)
 (esterification by, of (chlorophenoxy)ethanol)
 IT 96-49-1, Ethylene carbonate
 RL: RCT (Reactant)
 (etherification by, of chlorophenol)
 IT 25620-62-6, Dibromoethane

RL: RCT (Reactant)
 (etherification by, of trichlorophenol, in prepn. of agrochem.
 fungicide)
 IT 88-06-2, 2,4,6-Trichlorophenol
 RL: RCT (Reactant)
 (etherification of, by dibromoethane)
 IT 106-48-9, 4-Chlorophenol
 RL: RCT (Reactant)
 (etherification of, by ethylene carbonate)
 IT 26378-23-4P, 2-(2-Bromoethoxy)-1,3,5-trichlorobenzene
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and alkylation by, of propylamine, in prepn. of agrochem.
 fungicide)
 IT 67747-01-7P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and amidation by, of thiazole carboxylate, in prepn.
 of agrochem. fungicides)
 IT 14527-41-4P, 5-Thiazolecarboxylic acid
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and amidation of, in prepn. of agrochem. fungicide)
 IT 1892-43-9P, 2-(4-Chlorophenoxy)ethanol
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and esterification of, by chlorobenzenesulfonyl chloride)
 IT 14527-44-7P, Methyl 5-thiazolecarboxylate 41125-73-9P, 5-
 Thiazolecarbonyl chloride
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and sapon. of)
 IT 121046-39-7P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and N-alkylation by, of thiazolecarboxamides, in
 pregn. of agrochem. fungicides)
 IT 121046-38-6P, N-Propyl-5-thiazolecarboxamide
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pregn. and N-alkylation of, in prepn. of agrochem. fungicides)
 IT 121045-99-6P 121046-00-2P 121046-01-3P 121046-02-4P 121046-03-5P
 121046-04-6P 121046-05-7P 121046-06-8P 121046-07-9P 121046-08-0P
 121046-09-1P 121046-10-4P 121046-11-5P 121046-12-6P 121046-13-7P
 121046-14-8P 121046-15-9P 121046-16-0P 121046-17-1P 121046-18-2P
 121046-19-3P 121046-20-6P 121046-21-7P 121046-22-8P 121046-23-9P
 121046-24-0P 121046-25-1P 121046-26-2P 121046-27-3P 121046-28-4P
 121046-29-5P 121046-30-8P 121046-31-9P 121046-32-0P 121046-33-1P
 121046-34-2P 121046-35-3P 121046-36-4P 121046-37-5P 121055-23-0P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except
 adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP
 (Preparation); USES (Uses)
 (pregn. of, as agrochem. fungicide)

L7 ANSWER 33 OF 33 CAPLUS COPYRIGHT 1999 ACS
 AN 1989:231447 CAPLUS
 DN 110:231447
 TI Alpha-unsaturated amines, particularly 1,1-diamino-2-nitroethylene
 derivatives, their insecticidal/miticidal compositions, and processes for
 their preparation
 IN Minamida, Isao; Iwanaga, Koichi; Okauchi, Tetsuo
 PA Takeda Chemical Industries, Ltd., Japan
 SO Eur. Pat. Appl., 118 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 IC ICM C07D213-36
 ICS C07D213-74; C07D215-12; C07D277-38; C07D213-71; C07D277-28;
 C07D277-32; C07D213-61; A01N043-40; A01N043-42; A01N043-78
 CC 27-16 (Heterocyclic Compounds (One Hetero Atom))
 Section cross-reference(s): 5, 28
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 302389	A2	19890208	EP 1988-112210	19880728
	EP 302389	B1	19931222		
	EP 302389	A3	19900131		
	R: AT, BE, CH, DE, ES, FR, IN 167709	GB, GR, IT, LI, LU, NL, SE			
	EP 529680	A	19901208	IN 1988-MA493	19880712
	EP 529680	A2	19930303	EP 1992-115873	19880720
	EP 529680	A3	19930714		
	EP 529680	B1	19980513		
	R: AT, BE, CH, DE, ES, FR, EP 509559	GB, GR, IT, LI, LU, NL, SE			
	EP 509559	A2	19921021	EP 1992-111470	19880728
	EP 509559	A3	19930630		
	R: AT, BE, CH, DE, ES, FR, IL 87250	GB, GR, IT, LI, LU, NL, SE			
	AT 98955	A1	19930610	IL 1988-87250	19880728
	ES 2061569	E	19940115	AT 1988-112210	19880728
	IL 100688	T3	19941216	ES 1988-112210	19880728
	AT 166051	A1	19950831	IL 1988-100688	19880728
	US 5849768	E	19980515	AT 1992-115873	19880728
	HU 53909	A	19981215	US 1988-225367	19880728
HU 204496	A2	19901228	HU 1988-4040	19880729	
HU 205076	B	19920128			
CN 1031079	B	19920330	HU 1990-722	19880729	
CN 1027447	A	19890215	CN 1988-104801	19880801	
JP 02000171	A2	19900105	JP 1988-192383	19880801	
JP 07014916	B4	19950222			
US 5175301	A	19921229	US 1989-406515	19890913	
IN 170790	A	19920523	IN 1990-MA378	19900516	
US 5214152	A	19930525	US 1991-655072	19910214	
JP 05345760	A2	19931227	JP 1993-8114	19930121	
JP 05345761	A2	19931227	JP 1993-8115	19930121	
JP 05345774	A2	19931227	JP 1993-8116	19930121	
JP 07049424	B4	19950531			
CN 1091737	A	19940907	CN 1993-114205	19931105	
CN 1093083	A	19941005	CN 1993-114206	19931105	
CN 1036649	B	19971210			
JP 07206820	A2	19950808	JP 1994-254221	19940926	
JP 2551392	B2	19961106			
JP 07224036	A2	19950822	JP 1994-254222	19940926	
JP 2551393	B2	19961106			
US 5935981	A	19990810	US 1997-957749	19971024	
PRAI	JP 1987-192793	19870801			
	JP 1987-258856	19871013			
	JP 1988-16259	19880126			
	JP 1988-64885	19880317			
	IN 1988-MA493	19880712			
	EP 1988-112210	19880728			
	IL 1988-87250	19880728			
	US 1988-225367	19880728			
	US 1989-406515	19890913			
OS	MARPAT 110:231447				
GI					



II

AB Title compds. X1X2C:CR1NR2(CnH2n)A [I; 1 of X1 or X2 = electron-attracting group, other = H or electron-attracting group; R1 = group attached through

a N atom; R2 = H, group attached through a C, N, or O atom; n = 0-2; A = heterocyclyl; R1 = (un)substituted NH₂ when R2 = H] are prep'd. as insecticides and miticides. Aminolysis of (MeS)2C:CHNO₂ by Me₂NH in refluxing aq. EtOH gave Me₂N(MeS)C:CHNO₂, which underwent a 2nd aminolysis

by 6-chloro-3-pyridylmethylamine in refluxing EtOH to give [(chloropyridylmethyl)amino](dimethylamino)nitroethylene II. At 500 ppm (spray) on rice seedlings, II gave 100% mortality of larval Nilaparvata lugens in 7 days.

ST pyridylmethylaminonitroethylene prep'n insecticide miticide;
thiazolylmethylaminonitroethylene prep'n insecticide acaricide;
nitroethylene heterocyclymethylamino prep'n insecticide miticide

IT Acaricides

Insecticides
((heterocyclymethylamino)nitroethylenes)

IT 3000-75-7P, N-Ethyl-N-(3-pyridylmethyl)amine 3364-76-9P,
4-Chloromethylthiazole 4226-36-2P, 2-(3-Pyridyl)ethyl chloride hydrochloride 6971-44-4P, N-Methyl-N-(4-pyridylmethyl)amine

7032-20-4P

16273-55-5P 19690-13-2P 19730-13-3P,

N-Propyl-N-(3-pyridylmethyl)amine

20173-04-0P 20173-12-0P, N-Butyl-N-(3-pyridylmethyl)amine

21035-59-6P,

N-Methyl-N-(2-pyridylmethyl)amine 21543-49-7P 23879-54-1P

31982-51-1P 34107-46-5P, 6-Methyl-3-pyridylmethanol 37669-64-0P,

5-Bromo-3-pyridylmethanol 38663-85-3P, 2-Methoxyethyl isothiocyanate 39204-47-2P, 2-Chloromethylpyrazine 39620-02-5P, 5-Bromonicotinoyl chloride 42330-59-6P, 2-Chloro-3-pyridylmethanol 42506-12-7P

49609-84-9P, 2-Chloronicotinoyl chloride 52426-66-1P,

6-Methyl-3-pyridylmethyl chloride 55019-90-4P 59670-91-6P,

1,1-Dimethyl-2-(3-pyridylmethylidene)hydrazine 61771-67-3P,

2-Methoxy-5-methylaminopyridine 61832-41-5P 62658-90-6P,

2-Methylthio-3-pyridylmethanol 63326-08-9P 63361-56-8P,

N-Benzyl-N-(3-pyridylmethyl)amine 66171-50-4P 70258-18-3P

71718-88-2P, N-(3-Pyridylmethylidene)benzylamine 73335-64-5P,

N-(5-Bromo-3-pyridylmethyl)-N-methylamine 73781-91-6P, Methyl

6-chloronicotinate 82674-16-6P 89581-84-0P, 2-Chloro-3-pyridylmethyl chloride 97004-04-1P 97936-43-1P 101990-45-8P, 6-Bromo-3-

pyridylmethyl bromide 105827-74-5P, 6-Fluoro-3-pyridylmethyl bromide

109859-96-3P 120277-69-2P, 5-Bromo-3-pyridylmethyl chloride

120739-60-8P 120739-61-9P 120739-62-0P 120739-63-1P 120739-64-2P,

N-(3-Pyridylmethylidene)ethylamine 120739-65-3P, N-(3-

Pyridylmethylidene)-2-methoxyethylamine 120739-66-4P,

N-(3-Quinolylmethylidene)methylamine 120739-67-5P, N-(3-

Pyridylmethylidene)-n-propylamine 120739-68-6P 120739-69-7P,

N-Methyl-N-(3-quinolylmethyl)amine 120739-70-0P, 1,1-Dimethyl-2-(3-

pyridylmethyl)hydrazine 120739-71-1P, 2,2-Dichloro-3-pyridylmethylamine

120739-72-2P, N-(2,6-Dichloro-3-pyridylmethyl)phthalimide 120739-73-3P,

N-(2,6-Dichloro-3-pyridylmethyl)-N-methylamine 120739-74-4P

120739-75-5P 120739-76-6P 120739-77-7P,

N-(6-Chloro-3-pyridylmethyl)-N-

ethylamine 120739-78-8P 120739-79-9P 120739-80-2P 120739-81-3P

120739-82-4P 120739-83-5P,

N-(6-Chloro-3-pyridylmethyl)-N-isopropylamine

120739-84-6P, 2-Chloro-5-methylaminopyridine 120739-85-7P

120739-86-8P, N-(2,6-Dimethyl-4-pyridylmethyl)-N-methylamine

120739-87-9P, 2,6-Dimethyl-4-pyridylmethyl chloride 120739-88-0P,

N-(2-Chloro-3-pyridylmethyl)-N-methylamine 120739-90-4P 120739-91-5P

120739-92-6P, N-(2-Methylthio-3-pyridylmethyl)-N-methylamine

120739-93-7P 120739-94-8P 120739-95-9P 120739-96-0P 120739-97-1P

120739-98-2P 120739-99-3P 120740-00-3P 120740-01-4P 120740-02-5P,

N-Methyl-N-(6-methyl-3-pyridylmethyl)amine 120740-03-6P,

N-(6-Fluoro-3-pyridylmethyl)-N-methylamine 120740-04-7P,

N-(6-Bromo-3-pyridylmethyl)-N-methylamine 120740-05-8P,

N-(6-Bromo-3-pyridylmethyl)-N-ethylamine 120740-06-9P, N-(2-Chloro-5-

(prepn. of, as insecticide and miticide)

IT 57-06-7, Allyl isothiocyanate 57-14-7, 1,1-Dimethylhydrazine 74-88-4, reactions 74-89-5, Methylamine, reactions 75-04-7, Ethylamine, reactions 75-15-0, Carbon disulfide, reactions 75-52-5, Nitromethane, reactions 78-39-7, Ethyl orthoacetate 79-22-1, Methyl chloroformate 85-41-6, Phthalimide 107-10-8, n-Propylamine, reactions 109-08-0, 2-Methylpyrazine 109-85-3, 2-Methoxyethylamine 110-89-4, Piperidine, reactions 115-80-0, Triethyl orthopropionate 122-51-0, Ethyl orthoformate 124-40-3, Dimethylamine, reactions 124-41-4, Sodium methoxide 373-88-6, 2,2,2-Trifluoroethylamine hydrochloride 500-22-1, Pyridine-3-aldehyde 542-85-8, Ethyl isothiocyanate 556-61-6, Methyl isothiocyanate 592-82-5 593-56-6, O-Methylhydroxylamine hydrochloride 622-78-6, Benzyl isothiocyanate 624-83-9, Methyl isocyanate 626-35-7, Ethyl nitroacetate 628-30-8, n-Propyl isothiocyanate 693-95-8 872-85-5, Pyridine-4-aldehyde 1074-82-4, Potassium phthalimide 1121-60-4, Pyridine-2-aldehyde 1424-54-0, Methanesulfonyl isothiocyanate 2253-73-8, Isopropyl isothiocyanate 2258-42-6, Formic acetic anhydride 2369-19-9, 2-Fluoro-5-methylpyridine 2942-59-8, 2-Chloronicotinic acid 3430-14-6 3510-66-5 3731-52-0, 3-Pyridylmethylamine 5006-66-6, 6-Hydroxynicotinic acid 5350-93-6, 5-Amino-2-chloropyridine 5470-70-2, Methyl 6-methylnicotinate 6142-06-9, 2-Methylaminothiazole 6293-56-7, 2-(3-Pyridyl)ethanol 6628-77-9, 5-Amino-2-methoxypyridine 6638-79-5, N,O-Dimethylhydroxylamine hydrochloride 6959-48-4, 3-Pyridylmethyl chloride hydrochloride 7664-41-7, Ammonia, reactions 7803-57-8, Hydrazine hydrate 13623-94-4, 1,1-Bis(methylthio)-2-nitroethylene 13669-42-6, 3-Quinolinecarboxaldehyde 18088-01-2, 2,6-Dimethyl-4-pyridylmethanol 18364-47-1, 3-Methylaminopyridine 20826-04-4, 5-Bromonicotinic acid 23100-12-1 41789-37-1, 2,6-Dichloro-3-pyridylmethyl chloride 43083-12-1, Trimethyl orthobutyrate 61832-41-5, 1-Methylamino-1-methylthio-2-nitroethylene 74470-23-8 90875-71-1 101990-45-8, 6-Bromo-3-pyridylmethyl bromide 105827-91-6, 2-Chloro-5-chloromethylthiazole 112110-07-3, 3-Amino-5-trifluoromethylpyridine 120739-84-6 120740-65-0 120741-33-5
RL: RCT (Reactant)
(reaction of, in prepn. of insecticides and miticides)

thiazolylmethyl)-N-methylamine 120740-07-0P 120740-08-1P
 120740-09-2P, N-(2-Chloro-5-thiazolylmethyl)phthalimide
 120740-10-5P 120740-11-6P 120740-12-7P,
 N-(6-Chloro-3-pyridylmethyl)-N-(2,2,2-trifluoroethyl)amine 120740-13-8P 120740-14-9P 120740-15-0P
 120740-16-1P 120740-17-2P 120740-18-3P 120740-19-4P 120740-20-7P
 120740-21-8P 120740-22-9P 120740-23-0P 120740-24-1P 120740-25-2P
 120740-26-3P 120740-27-4P 120740-28-5P 120740-29-6P 120740-30-9P
 120740-31-0P 120740-32-1P 120740-33-2P 120740-34-3P 120740-35-4P
 120740-36-5P 120740-37-6P 120740-38-7P 120740-39-8P 120740-40-1P
 120740-41-2P 120740-42-3P 120740-43-4P 120740-44-5P 120740-45-6P
 120740-46-7P 120740-47-8P 120740-48-9P 120740-49-0P 120740-50-3P
 120740-51-4P 120740-52-5P 120740-53-6P 120740-54-7P 120740-55-8P
 120740-56-9P 120740-57-0P 120740-58-1P 120740-59-2P 120740-60-5P
 120740-61-6P 120740-62-7P 120740-63-8P 120740-64-9P 120740-65-0P
 120740-66-1P 120740-67-2P 120740-68-3P 120740-69-4P 120740-70-7P
 120740-71-8P 120740-72-9P 120740-73-0P 120740-74-1P 120740-75-2P
 120740-76-3P 120740-77-4P 120740-78-5P 120740-79-6P 120740-80-9P
 120740-81-0P 120740-82-1P 120740-83-2P 120740-84-3P 120740-85-4P
 120740-86-5P 120740-87-6P 120740-88-7P 120740-89-8P 120740-90-1P
 120740-91-2P 120740-92-3P 120740-93-4P 120740-94-5P 120740-95-6P
 120740-96-7P 120740-97-8P 120740-98-9P 120740-99-0P 120741-00-6P
 120741-01-7P 120741-02-8P 120741-03-9P 120741-04-0P 120741-05-1P
 120741-06-2P 120741-07-3P 120741-08-4P 120741-09-5P 120741-10-8P
 120741-11-9P 120741-12-0P 120741-13-1P 120741-14-2P 120741-15-3P
 120741-16-4P 120741-17-5P, 6-Chloro-3-pyridylmethyl isothiocyanate
 120741-18-6P 120741-19-7P 120741-20-0P 120741-21-1P 120741-22-2P
 120741-23-3P 120741-24-4P, 6-Bromo-3-pyridylmethyl isothiocyanate
 120741-25-5P 120741-26-6P 120741-27-7P 120741-28-8P 120741-29-9P
 120741-30-2P 120741-31-3P 120741-32-4P 120770-88-9P 120770-89-0P
 120770-90-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. and reaction of, in prepn. of insecticides and miticides)

IT 120741-34-6P 120741-35-7P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of)

IT 90787-25-0P 120738-33-2P 120738-34-3P 120738-35-4P 120738-36-5P
 120738-37-6P 120738-38-7P 120738-39-8P 120738-40-1P 120738-41-2P
 120738-42-3P 120738-43-4P 120738-44-5P 120738-45-6P 120738-46-7P
 120738-47-8P 120738-48-9P 120738-49-0P 120738-50-3P 120738-51-4P
 120738-52-5P 120738-53-6P 120738-54-7P 120738-55-8P 120738-56-9P
 120738-57-0P 120738-58-1P 120738-59-2P 120738-60-5P 120738-61-6P
 120738-62-7P 120738-63-8P 120738-64-9P 120738-65-0P 120738-66-1P
 120738-67-2P 120738-68-3P 120738-69-4P 120738-70-7P 120738-71-8P
 120738-72-9P 120738-73-0P 120738-74-1P 120738-75-2P 120738-76-3P
 120738-77-4P 120738-78-5P 120738-79-6P 120738-80-9P 120738-81-0P
 120738-82-1P 120738-83-2P 120738-84-3P 120738-85-4P 120738-86-5P
 120738-87-6P 120738-88-7P 120738-89-8P 120738-90-1P 120738-91-2P
 120738-92-3P 120738-93-4P 120738-94-5P 120738-95-6P 120738-96-7P
 120738-97-8P 120738-98-9P 120738-99-0P 120739-00-6P 120739-01-7P
 120739-02-8P 120739-03-9P 120739-04-0P 120739-05-1P 120739-06-2P
 120739-07-3P 120739-08-4P 120739-09-5P 120739-10-8P 120739-11-9P
 120739-12-0P 120739-13-1P 120739-14-2P 120739-15-3P 120739-16-4P
 120739-17-5P 120739-18-6P 120739-19-7P 120739-20-0P 120739-21-1P
 120739-22-2P 120739-23-3P 120739-24-4P 120739-25-5P 120739-26-6P
 120739-27-7P 120739-28-8P 120739-29-9P 120739-30-2P 120739-31-3P
 120739-32-4P 120739-33-5P 120739-34-6P 120739-35-7P 120739-36-8P
 120739-37-9P 120739-38-0P 120739-39-1P 120739-40-4P 120739-41-5P
 120739-42-6P 120739-43-7P 120739-44-8P 120739-45-9P 120739-46-0P
 120739-47-1P 120739-48-2P 120739-49-3P 120739-50-6P 120739-51-7P
 120739-52-8P 120739-53-9P 120739-54-0P 120739-55-1P 120739-56-2P
 120739-57-3P 120739-58-4P 120739-59-5P 120741-36-8P,
 1,1-Bis[(3-pyridylmethyl)amino]-2-nitroethylene 120741-37-9P
 120770-85-6P 120770-86-7P 120770-87-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)

=> s 13

L4 64 L3

=> s 14 and process

L5 1206643 PROCESS
4 L4 AND PROCESS

=> s 14 and halogen?

L6 93976 HALOGEN?
0 L4 AND HALOGEN?

=> d 15 1-4 ibib ab hitstr

L5 ANSWER 1 OF 4 CAPLUS COPYRIGHT 1999 ACS
ACCESSION NUMBER: 1999:609863 CAPLUS
DOCUMENT NUMBER: 131:253657
TITLE: Discovery of a new systemic insecticide, nitenpyram
and its insecticidal properties
AUTHOR(S): Akayama, Atsuo; Minamida, Isao
CORPORATE SOURCE: Agricultural Research Laboratories, Agro Company,
Takeda Chemical Industries, Ltd., Tsukuba, 300-42,
Japan
SOURCE: Nicotinoid Insectic. Nicotinic Acetylcholine Recept.,
[Symp.] (1999), Meeting Date 1997, 127-148.
Editor(s): Yamamoto, Izuru; Casida, John E.

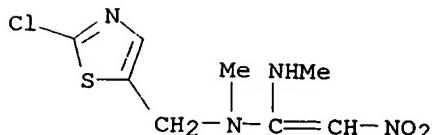
Springer:
Tokyo, Japan.
CODEN: 68EFAV
DOCUMENT TYPE: Conference
LANGUAGE: English
AB A large no. of nitromethylene heterocyclic compds. were screened against
Nilaparvata lugens and the Nephrotettix cincticeps. Nitenpyram was
discovered in the process of the optimization of the
substituents of an acyclic nitroethene. The compd. maintains poor
photostability; however, it shows excellent residual effect against rice
hoppers and aphids by foliar spray. The poor photostability of
nitenpyram
is considered to be advantageous for environmental safety and for
minimization of the resurgence of the phytophagous mite and adverse
effects
against beneficial insects. Nitenpyram is highly sol. in water, and
shows

excellent systemic action and no phytotoxicity, which characteristics
enable the various application methods of the compd. The compd. has been
developed for controlling the brown planthopper, however, it is highly
active against some other pest species, and single soil application of
nitenpyram effectively controls aphids, whiteflies, trips, and the
serpentine leaf miner, which are the major greenhouse pests difficult to
control by conventional insecticides.

IT 120739-30-2 120739-31-3
RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
(insecticidal activity against Nilaparvata lugens and Nephrotettix
cincticeps)

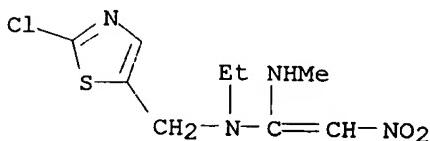
RN 120739-30-2 CAPLUS

CN 1,1-Ethenediamine,
N-[(2-chloro-5-thiazolyl)methyl]-N,N'-dimethyl-2-nitro-



RN 120739-31-3 CAPLUS

CN 1,1-Ethenediamine, N-[(2-chloro-5-thiazolyl)methyl]-N-ethyl-N'-methyl-2-nitro- (9CI) (CA INDEX NAME)



L5 ANSWER 2 OF 4 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1998:668115 CAPLUS

DOCUMENT NUMBER: 129:290052

TITLE:

Process for the preparation of nitroguanidine derivatives starting from 2-nitroiminohexahydro-1,3,5-triazines in the presence of ammonia, primary or secondary amines

INVENTOR(S): Ebihara, Koichi; Ura, Daisuke; Miyamoto, Michihiko; Kaiho, Tatsuo

PATENT ASSIGNEE(S): Mitsui Chemicals, Inc., Japan

SOURCE: Eur. Pat. Appl., 13 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 869120	A1	19981007	EP 1998-105850	19980331
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
CN 1197064	A	19981028	CN 1998-108267	19980331
JP 11236381	A	19990831	JP 1998-86842	19980331
PRIORITY APPLN. INFO.:			JP 1997-80178	19970331
			JP 1997-82838	19970401
			JP 1997-223813	19970820
			JP 1997-258968	19970924
			JP 1997-347934	19971217

OTHER SOURCE(S): MARPAT 129:290052

AB Described is a **process**, as a substitute for hydrolysis, for prep. a nitroguanidine deriv., RCH₂NHC(:NNO₂)NHR₂ (R = 2-chloro-5-pyridyl, 2-chloro-5-thiazolyl, 2-, 3-tetrahydrofuryl, 2-methyl-4-tetrahydrofuryl; R₂ = Me, allyl), which comprises treating a triazine, I (R₁ = Me, benzyl, i-Pr, Et, t-Bu, cyclohexyl; R, R₂ = same as above), with NH₃, a primary amine or a secondary amine, or a salt thereof.

IT 131748-59-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 131748-59-9 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitro- (9CI)